



FOIA Documents prove COVID-19 "Vaccines" were treated as "Medical Counter-measures" to a Bio-Weapon attack. Entire COVID Operation Run by US National Security Council with NO MANUFACTURING SAFEGUARDS OR DISCLOSURES

[NATION HAL TURNER](#) 08 JANUARY 2023 HITS: 206

New Docs Reveal Department of Defense Controlled COVID-19 Program from the start. The FDA Vaccine Approval Process was Theater. A combination of the PREP Act, Emergency Use Authorization, and Other Transactions Authority (OTA), shielded Big Pharma, Agencies, and Medical Participants that Delivered Unregulated Vaccines, from any liability.

According to congressionally passed statutes, research of active laws, and extra details obtained through the Freedom of Information Act, the Department of Defense owns, implements, and oversees the COVID-19 vaccine program as a "Countermeasure" to foreign attack. While the public was bombarded with an orchestrated fear campaign, the U.S. Government managed the Covid response as a national security threat.

[The research and documents](#) were obtained by a former executive of a pharmaceutical Contract Research Organization (CRO), Sasha Latypova, [and intensive legal researcher](#) Katherine Watt.

The Three-Legged Stool

The undercover operation was orchestrated utilizing three critical legal maneuvers:

1. Emergency Use Authorization EUA.
2. Prep Act,
3. Other Transactions Authority

President Trump declared a Public [Health](#) Emergency (PHE) on March 13, 2020, under the Stafford Act, putting the National Security Council in charge of the Covid policy. Covid-19 vaccines are "**medical countermeasures**" – a grey area of products that are not regulated as vaccines or medicines.

"They put the National Security Council in charge and treated it as an act of war," said Latypova.

According to Operation Warp Speed/ASPR reports, the DoD ordered, oversaw, and tightly managed the development, manufacture, and distribution of Covid countermeasures, mainly

utilizing the DoD's previously established network of military contractors and consortia.

Department of Defense (DoD), the *Biomedical Advanced Research and Development Authority* (BARDA), and Department of [Health](#) and Human Services (HHS), ordered all Covid countermeasures, including "vaccines" as prototype demonstrations of large-scale manufacturing, avoiding regulations and transparency under Other Transaction Authority. As prototypes used under EUA during PHE, Covid countermeasures, including "vaccines," **need not comply with the U.S. laws for manufacturing quality, safety, and labeling.**

"The implication is that the U.S. Government authorized and funded the deployment of noncompliant biological materials on Americans without clarifying their "prototype" legal status, making the materials not subject to normal regulatory oversight, all while maintaining a fraudulent pseudo- "regulatory" presentation to the public," said Latypova.

"Most incredible is the fact that current Laws enacted by the United States Congress appear to make the coverup actions LEGAL!"

Under the PHE, medical countermeasures are not regulated or safeguarded as pharmaceutical products (21 USC 360bbb-3(k)).

The American people were led to believe that the FDA, CDC, and figureheads like Anthony Fauci, oversaw the COVID-19 vaccine program. Their involvement was an orchestrated information operation. All decisions concerning the COVID-19 vaccine research, materials acquisition, distribution, and information sharing were tightly controlled by the DoD.

Hundreds of Covid countermeasures [contracts](#) have been uncovered. Many disclosures are in redacted form. However, Latypova and Watt have found sources to fill in the details.

A review of these contracts indicates a high degree of control by the U.S. Government (DoD/BARDA). It specifies the scope of deliverables as "demonstrations" and "prototypes" only while excluding clinical trials and manufacturing quality control from the scope of work paid for by the contracts.

To ensure that the Pharma is free to conduct the fake clinical trials without financial risk, the contracts include the removal of all liability for the manufacturers and any contractors along the supply and distribution chain under the 2005 PREP Act and related federal legislation.

Why is no action by regulators or courts? According to Latypova and Watt, a combination of recently passed legislation and executive orders make it **LEGAL to LIE!** The HHS Secretary is accountable to no one if the [Health](#) National Emergency continues to be extended by Congress every three months.

A significant information operation was set in motion the minute COVID-19 hit. The U.S. government, the intelligence community, the media, and Big Tech colluded to orchestrate and implement an intense pressure campaign designed to get the vaccine legally designated under the Emergency Use Authorization Act while vilifying dissenting doctors, critics, and viable alternative treatments. This designation allowed for speedy manufacturing devoid of the standard safety and public [health](#) protocols.

For a vaccine to receive designation under the EUA, there can be no other known treatments or cures. Therefore, many proven treatments such as ivermectin and hydroxychloroquine were blacklisted in the media and dismissed as “[horse](#) de-wormers” when these cheap, readily available drugs were in the past heralded for their effectiveness.

Eminent COVID-treating doctors such as Peter M. McCullough and Pierre Kory have faced unprecedented attacks on their medical credentials.

<https://haltturnerradioshow.com/index.php/en/news-page/news-nation/foia-documents-prove-covid-19-vaccines-were-treated-as-medical-counter-measures-to-a-bio-weapon-attack-entire-covid-operation-run-by-us-national-security-council-with-no-manufacturing-safeguards-or-disclosures>

The role of the US DoD (and their co-investors) in "covid countermeasures" enterprise.

Intelligence is never artificial.



[Sasha Latypova](#)

Dec 28, 2022

424

325

I asked the AI Chat Bot “What was the role of the DoD in covid-countermeasures?” Here is what it wrote:

“The Department of Defense (DoD) played a significant role in the development of countermeasures against COVID-19, the disease caused by the novel coronavirus. The DoD worked closely with other federal agencies, such as the Department of [Health](#) and Human Services (HHS), to respond to the pandemic. One of the main roles of the DoD in the COVID-19 response was to provide logistical support and medical resources to help healthcare workers and first responders on the front lines of the pandemic. The DoD also worked to develop

and distribute personal protective equipment (PPE) and other supplies to support the response efforts.

In addition, the DoD provided funding and resources to support the development of vaccines and treatments for COVID-19. The Defense Advanced Research Projects Agency (DARPA), a research arm of the DoD, played a key role in this effort by funding and supporting the development of treatments and vaccines through its "Rapid Response Research" program.

Overall, the DoD played a vital role in the response to the COVID-19 pandemic by providing logistical support, medical resources, and funding for research and development efforts to combat the disease.”

AI Chat Bot gets a “F” - for a highly controlled narrative answer, which was fed to all MSM sources, while the “alternative” media did not examine it very much, with some exceptions. It wasn’t just logistics or even just lots of funding, and certainly nobody was “combatting disease”.

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The truth is much more interesting than fiction:

All Covid countermeasures, including the biological warfare agents marketed as “Covid-19 vaccines”, were ordered by the US DoD as a “large scale manufacturing demonstration” via Other Transactions Authority contracts.

Hundreds of Covid countermeasures [contracts](#) became available via FOIA and SEC disclosures in redacted form. Review of these contracts indicates a high degree of control by the US Government (DoD/BARDA) and specifies the scope of deliverables as “demonstrations” and “prototypes” only. In other words, the US Government and DOD specifically ordered a fake theatrical performance from the pharmaceutical manufacturers. Just to make extra certain that the pharmas are free to conduct the fakery, the contracts include the removal of all liability for the manufacturers and any contractors along the supply and distribution chain under the 2005 PREP Act and related federal legislation.

The contracts are structured under Other Transactions Authority (OTA) - [OTA](#) method of contracting allows federal agencies to [order](#) otherwise-regulated products bypassing any such regulations, as well as financial accountability mechanisms that cover standard government

contracting, and other laws that regulate disclosure and Intellectual Property (IP) derived from publicly funded research.

“Other” is a catchall category that is not a contract, not a research grant, not a procurement, etc.: not any normally regulated/accountable government contracting.

Here is a typical contract scope for “vaccines”:



**DEPARTMENT OF THE ARMY
U.S. ARMY CONTRACTING COMMAND – NEW JERSEY
PICATINNY ARSENAL, NEW JERSEY 07806-5000**

REPLY TO
ATTENTION OF

21 July 2020

Army Contracting Command – New Jersey
ACC-NJ, Building 9
Picatinny Arsenal, NJ 07806

SUBJECT: Technical Direction Letter for Medical CRBN Defense Consortium (MCDC), Request for Prototype Proposals (RPP) 20-11, Objective PRE-20-11 for “COVID-19 Pandemic – Large Scale Vaccine Manufacturing Demonstration” (Pfizer, Inc.)

REF: Prizer Request for Technical Direction Letter, RPP 20-11 under OTA W15QKN-16-9-1002 for Objective PRE-20-11, dated 20 July 2020

Advanced Technology International
ATTN: (b) (6), Sr. Contracts Manager
315 Sigma Drive
Summerville, SC 29486

Dear (b) (6),

The Army Contracting Command – New Jersey (ACC-NJ), in supporting the Joint Project Manager – Medical Countermeasure Systems (JPM-MCS), issued MCDC RPP 20-11 on 09 June 2020. Members of the MCDC submitted proposals in accordance with this RPP. The Government received and evaluated all proposal(s) submitted and a Basis of Selection has been executed, selecting Pfizer, Inc. as the awardee. The Government requests that a Firm-Fixed-Price Project Agreement be issued to Pfizer, Inc. to award this proposal under Other Transaction Agreement W15QKN-16-9-1002, to be performed in accordance with the attached Government Statement of Work (SOW).

Based upon the acceptable update of Pfizer, Inc.’s proposal for “COVID-19 Pandemic – Large Scale Vaccine Manufacturing Demonstration” and 1) The Project Agreement Recipient’s concurrence with the requirements included in the Government SOW; 2) An acceptable milestone schedule that meets SOW requirements, and; 3) The price proposed that has been analyzed by the Government, you are hereby directed to issue a Project Agreement to Pfizer, Inc. for the subject project. The total project value has been determined fair and reasonable and Pfizer, Inc.’s proposal has been selected IAW the above referenced Basis of Selection.

The total approved cost to the Government for this effort is not to exceed \$1,950,097,500.00. The break-out of the costs is as follows: \$1,950,000,000.00 to perform project efforts included in the SOW and \$97,500.00 for the Consortium Management Firm (CMF) Administrative Cost. The CMF Administrative Cost was approved as a “Special Allocation” for Operation Warp Speed (OWS) Prototype Projects executed under the MCDC OTA. The effort currently has \$1,950,097,500.00 of available funding, comprised of \$1,950,000,000.00 for the Project Agreement, \$67,500.00 for the

While the DOD/BARDA countermeasure contracts refer to safety and efficacy requirements for vaccines and mention current Good Manufacturing Practices (cGMP) compliance, these items are explicitly carved out as not being paid for nor ordered by the US Government.

1.2 Scope

The scope of this prototype project is the demonstration by Pfizer of the supply and logistics capability to manufacture and distribute to the Government of 100M doses of a novel mRNA-based vaccine that has received FDA-approval or authorization based on demonstration of efficacy (hereafter FDA-approved or authorized). The criteria for successful Emergency Use Authorization (EUA) are described in *Emergency Use Authorization of Medical Products and Related Authorities: Guidance for Industry and Other Stakeholders, January 2017*; and *Development and Licensure of Vaccine to Prevent COVID-19: Guidance for Industry June 2020*. The successful provision of these doses shall establish the effectiveness of a technology capable of potentially providing immediate and long-term solutions to coronavirus infections. While pre-clinical, clinical, and chemistry/manufacturing/controls (CMC) activities are described in the Background section of this Statement of Work, the Parties acknowledge and agree that such activities not related to the large-scale manufacturing demonstration are out-of-scope for this prototype project as Pfizer and BioNTech have and will continue to fund these activities, without the use of Government funding.

8

This Statement of Work includes proprietary and confidential commercial data of Pfizer Inc. that shall not be disclosed outside the MCDC Management Firm and the Government and shall not be duplicated, used, or disclosed, in whole or in part, for any purpose other than to evaluate this Statement of Work and negotiate any subsequent award. If, however, an agreement is awarded as a result of, or in connection with, the submission of this data, the MCDC Management Firm and the Government shall have the right to duplicate, use, or disclose these data to the extent provided in the resulting agreement. This restriction does not limit the MCDC Management Firm and the Government's right to use the information contained in these data if they are obtained from another source without restriction. The data subject to this restriction are set forth on each page of this Statement of Work.

US 168054648v17

This gets even more interesting when we examine some of the redactions in contracts:

1.1.1 BACKGROUND

(b) (4), Pfizer and BioNTech entered into an agreement for the co-development and distribution (excluding China) of a potential mRNA-based coronavirus vaccine aimed at preventing COVID-19 infection (the “Pfizer/BioNTech Agreement”). Under the Pfizer/BioNTech Agreement, (b) (4)

for the prevention of COVID-19 (b) (4)

for the prevention of COVID-19 (b) (4)

(b) (4)

2

This Statement of Work includes proprietary and confidential commercial data of Pfizer Inc. that shall not be disclosed outside the MCDC Management Firm and the Government and shall not be duplicated, used, or disclosed, in whole or in part, for any purpose other than to evaluate this Statement of Work and negotiate any subsequent award. If, however, an agreement is awarded as a result of, or in connection with, the submission of this data, the MCDC Management Firm and the Government shall have the right to duplicate, use, or disclose these data to the extent provided in the resulting agreement. This restriction does not limit the MCDC Management Firm and the Government's right to use the information contained in these data if they are obtained from another source without restriction. The data subject to this restriction are set forth on each page of this Statement of Work.

US 168054648v17

I know what is in the redacted part of the above paragraph and it was not hard to figure out. The first redaction under 1.1.1 BACKGROUND is “Fosun Pharmaceuticals”, so the sentence reads “Fosun Pharmaceuticals”, Pfizer and BioNTech entered into an agreement for the co-development...”

Note: the only journalist I am aware of in either “mainstream” or “resistance” who mentioned Fosun was Naomi [Wolf](#), kudos to her. I was in touch with The Epoch Times to try to publish this information, and even they decided to bury the story (but they published my other materials). I did discuss this on Dr. Jane Ruby’s show, and kudos to her as well for not being afraid to cover the truth.



BioNTech and Fosun Pharma form COVID-19 vaccine strategic alliance in China

March 16, 2020

- *BioNTech and Fosun Pharma will jointly conduct clinical trials of BNT162 in China, leveraging BioNTech's proprietary mRNA vaccine technology and Fosun Pharma's clinical development and commercialization capabilities in China*
- *Fosun Pharma will commercialize the vaccine in China upon regulatory approval, with BioNTech retaining full rights to develop and commercialize the vaccine in the rest of the world*
- *Fosun Pharma will pay BioNTech up to USD 135M (EUR 120M) in upfront and potential future investment and milestone payments; the two companies will share future gross profits from the sale of the vaccine in China*

MAINZ, Germany, and SHANGHAI, China, March 16, 2020 (GLOBE NEWSWIRE) -- [BioNTech SE](#) (Nasdaq: BNTX, "BioNTech") and [Shanghai Fosun Pharmaceutical \(Group\) Co., Ltd](#) ("Fosun Pharma" or "Group"; Stock Symbol: 600196.SH, 02196.HK) announced today a strategic development and commercialization collaboration to advance BioNTech's mRNA vaccine candidate BNT162 in China for the prevention of COVID-19 infections.

Under the terms of the agreement, the two companies will work jointly on the development of BNT162 in China. The companies will collaborate to conduct clinical trials in China leveraging Fosun Pharma's extensive clinical development, regulatory, and commercial capabilities in the country.

If approved, Fosun Pharma will commercialize the vaccine in China. BioNTech will supply the mRNA vaccine for clinical trials from GMP manufacturing facilities in Europe along with its partner [Polymun](#). BioNTech will retain full rights to develop and commercialize the vaccine in the rest of the world.

Pfizer-BioNTech is really a 3-party R&D alliance: Fosun-Pfizer-BioNTech, and by "party" I mean that one of the three is the Chinese Communist Party. Fosun is a huge Chinese [conglomerate](#) that owns a large number of global companies, and its chairman [Guo Guangchang](#) is a very high ranking member of the CCP. It is curious that the US DoD awarded \$10 billion (Pfizer's Operation Warp Speed/DoD/BARDA contract) to a venture whose substantial equity (and IP) holder is the the Chinese Communist Party. For avoidance of doubt:

CONTINUATION SHEET	Reference No. of Document Being Continued PHIN/SHIN W15QKN-21-C-0012 MOD/AMD	Page 2 of 51
Name of Offeror or Contractor: PFIZER INC.		

SUPPLEMENTAL INFORMATION

Buyer Name: (b) (6)
Buyer Office Symbol/Telephone Number: CCNJ-IC/(b) (6)
Type of Contract 1: Firm Fixed Price
Kind of Contract: Supply Contracts and Priced Orders
Type of Business: Large Business Performing in U.S.
Surveillance Criticality Designator: A

*** End of Narrative A0000 ***

Executive Summary

Background:

The Department of Health and Human Services (DHHS) continuously monitors emerging infectious disease risk and prepares to respond to the threat of novel emerging infectious disease outbreaks in the United States. DHHS is responding to an outbreak of respiratory disease caused by a novel coronavirus that was first detected in China and which has now spread to worldwide, including in the United States. The virus has been named SARS-CoV-2 and the disease it causes has been named coronavirus disease 2019 (abbreviated COVID-19).

On January 30, 2020, the International Health Regulations Emergency Committee of the World Health Organization (WHO) declared the outbreak a public health emergency of international concern (PHEIC). On January 31, Health and Human Services Secretary Alex M. Azar II declared a public health emergency (PHE) for the United States to aid the nations healthcare community in responding to COVID-19. On March 11, WHO publicly characterized COVID-19 as a pandemic. On March 13, the President of the United States declared the COVID-19 outbreak a national emergency. Vaccination is often the most effective measure for the control of infectious diseases. In July 2020, the DOD awarded an Other Transaction Agreement under the authority 10 USC 2371b to Pfizer to scale up manufacture of their BNT162b2 vaccine candidate. The candidate entered Phase 3 clinical trials and scale up of their manufacturing processes. On November 9, 2020, Pfizer announced that BNT162b2 was >90% effective based on interim analysis of partial data from their Phase 3 clinical trial. On November 18, 2020, Pfizer reported 95% effectiveness based on analysis of a larger dataset that included 170 confirmed cases among the Phase 3 volunteers (162 in the placebo group and 8 in the vaccinated group). Based on the strength of this data, Pfizer formally requested Emergency Use Authorization (EUA) from the US Food and Drug Administration.

1. This action has a total Firm Fixed Price value of \$10,016,418,500 inclusive of all options. At this time, CLINs 0001, 0002, and 0003 are funded in the amount of \$2,011,282,500.

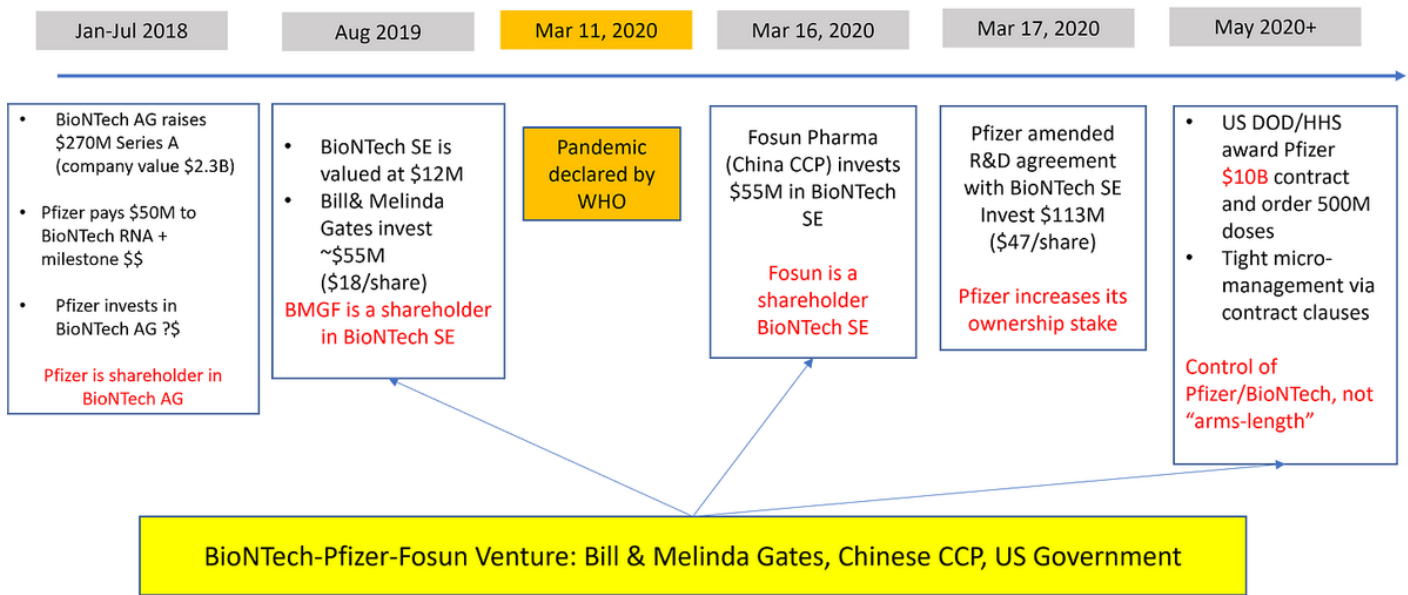
2. The Representations and Certifications made by Pfizer in the System for Award Management (SAM) are hereby incorporated into this contract by reference.

3. The Pfizer Small Business Subcontracting Plan, dated 01 March 2020 is hereby incorporated into the contract (see Attachment 0002).

*** END OF NARRATIVE A0001 ***

Below is the timeline of some of the key investments and R&D deals I was able to identify from public SEC shareholder disclosures, immediately preceding and following the “pandemic”:

Timeline of Select Contracts



Just to make sure, we are talking about the exact technology in the mRNA shots. Here is the definition from March 17, 2020 agreement between Pfizer and BioNTech (p. 4):

whether pending or issued that (a) is Controlled by BioNTech or any of its Affiliates as of the Effective Date or comes into the Control of BioNTech or any of its Affiliates during the Term (other than, in either case, through the grant of a license by Pfizer) and (b) claims any BioNTech Know-How.

1.17 **"BioNTech Technology"** means the BioNTech Patent Rights, BioNTech Materials, BioNTech Know-How. For avoidance of doubt, BioNTech Technology includes all Intellectual Property Rights Controlled by BioNTech pursuant to the Fosun Agreement.

The same document describes a data sharing agreement, "pharmacovigilance" globally among the 3 parties. They will count the bodies and share the data with each other:

associated with Candidates or Products. BioNTech shall be responsible for maintaining a suitable safety database.

8.3.3 The Parties acknowledge and agree that they have entered into a pharmacovigilance agreement covering pharmacovigilance responsibility relating to Development Activities and shall update such agreement or enter into a new pharmacovigilance agreement with respect to Commercialization Activities (each a “Pharmacovigilance Agreement”), in each case reflecting the applicable terms set forth in Section 8.3.7 and Schedule 8.3.

8.3.4 Following the filing of the IND for Candidate(s) with FDA:

8.3.4.1 should BioNTech require Pfizer to take over certain activities in relation to collecting, monitoring, evaluating, sharing and reporting to applicable Governmental Authorities, but excluding Ethics Committees, information regarding patient safety (including adverse drug) experiences that are or may be associated with Candidates or Products in the EU, the Parties shall agree and execute an amendment to the Pharmacovigilance Agreement to (i) reflect the additional activities and responsibilities the Parties have agreed Pfizer will perform in the EU, and (ii) set out the procedures the Parties have agreed upon to allow for the reconciliation of BioNTech’s safety database with Pfizer’s safety database. The effectiveness of the amendment shall be conditional upon BioNTech delivering to Pfizer (x) confirmation from the relevant Governmental Authorities in the EU that they have accepted an amendment to the clinical trial protocol for any on-going clinical trial of Candidates or Product in the EU to reflect the necessary changes (as agreed with Pfizer) in responsibilities and contact information for collecting, monitoring, evaluating, sharing and reporting of information regarding patient safety (including adverse drug) experiences, and (y) written confirmation from BioNTech that it has amended the relevant clinical trial agreements to reflect the change in pharmacovigilance provider and trained the investigators on the new reporting procedures; and,

8.3.4.2 BioNTech through their agreement with Fosun shall ensure that Fosun, via BioNTech, deliver to Pfizer (x) a copy of a due diligence report on Fosun’s safety data reporting system reasonably acceptable to Pfizer in terms of findings made, (y) a copy of the pharmacovigilance agreement between BioNTech and Fosun which, inter alia, provides for delivery to Pfizer of fully assessed, translated (into English) CIOMS forms for all SAEs: Death / life threatening SUSARs – 5 Business Days from Day 0 (Day 0 being receipt by Fosun from the clinical investigator), or 10 days for all other SAEs, [***] and (z) details of the quality management system used with Fosun to ensure that if late inbound reports are received BioNTech can request root cause analysis and implementation of corrective and preventive actions by Fosun. The Parties agree that prior to Fosun’s commencement of clinical activities by Fosun, BioNTech shall have entered into a written agreement with Fosun, reflecting the foregoing.

8.3.5 The Pharmacovigilance Agreement and each amendment to it from time to time shall set forth the responsibilities and procedures for (i) collecting, monitoring, evaluating, sharing and reporting to applicable Governmental Authorities information regarding patient safety (including adverse drug) experiences that are or may be associated with Candidates or Products in the countries covered by that agreement and (ii) providing regulatory information to and support

On the “pharmacovigilance” aspect, there is a 4th participant in this arrangement - the Israeli Ministry of [Health](#), which entered into a data sharing [agreement](#) with Pfizer on January 6, 2021 and gave Pfizer (and by extension, US DoD and anyone who controls it, BioNTech and anyone who controls it, Fosun and anyone who controls it, i.e. CCP) access to all their citizens’ centralized electronic [health](#) records. But don’t worry, Benjamin Netanyahu promised to keep the data de-identified. Right.

REAL-WORLD EPIDEMIOLOGICAL EVIDENCE COLLABORATION AGREEMENT

This REAL-WORLD EPIDEMIOLOGICAL EVIDENCE COLLABORATION AGREEMENT dated as of January 6, 2021 (this “**Agreement**”) by and between the Israeli Ministry of Health, acting on its own behalf and on behalf of the State of Israel (the “**MoH**”), and Pfizer Inc., a Delaware corporation (together with its Affiliates, “**PFIZER**”) (each, a “**Party**” and, collectively, the “**Parties**”).

WHEREAS, PFIZER and BioNTech SE, a company organized and existing under the laws of Germany are collaborating to develop a vaccine to address the global COVID-19 pandemic; and

WHEREAS, the Parties had previously entered into the confidential Manufacturing and Supply Agreement dated [REDACTED] (the “**Manufacturing and Supply Agreement**”), under which MoH agreed to purchase the Product (as defined below) and PFIZER agreed to manufacture and supply the Product, all in accordance with the terms of the Manufacturing and Supply Agreement, and subject to certain conditions precedent, including but not limited to certain regulatory approvals and supply availability; and

WHEREAS, under Section 2.1(f) of the Manufacturing and Supply agreement, the Parties agreed to cooperate on a reasonable basis to share information and data regarding the distribution, administration and use of the Product, including to track its benefits; and

WHEREAS, PFIZER has obtained certain conditional approvals for the Product, including under Regulation 29(a)(9) of the Israeli *Pharmacist Regulations (Medical Preparations)*, 1986, as amended, and analogous emergency use authorizations in other jurisdictions; and

WHEREAS, the Parties agree that it would be highly beneficial from a public health perspective to track pandemic data in accordance with vaccination compliance in a Real-World context to evaluate whether herd immunity protection is observed during the Product vaccination program rollout.

NOW THEREFORE, for and in consideration of the premises and mutual covenants and agreements contained herein and for other good and valuable consideration, the receipt and sufficiency of which are hereby acknowledged, the Parties hereto intending to be legally bound, hereby agree as follows:

1. DEFINITIONS

Side note - Israeli government recently “misplaced” the Manufacturing and Supply Agreement with Pfizer mentioned in the data sharing agreement above (so we know for sure it exists). The government sadly cannot find it for some reason...



bad cattitude

israeli government "lost the agreement" with pfizer for covid vaccines

so, apparently, the great risk of working from home is the VAST seeming increase in “the dog eating your homework.” but this is really something. israel, who everyone around the world was looking to fo...

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This gets even larger and more interesting when looking at the sources of “R&D” financing. Turns out, there were numerous financial backers and co-investors in the BioNTech “venture” in the years preceding the global fraud and mass murder exercise. According to Crunchbase,

BioNTech, a tiny company with just a handful of employees and NO PRODUCTS or scale manufacturing, raised \$1.7B in 9 rounds of investments since around 2008. Large portion of the money, \$1B+ was raised before 2020. What was it for, since no big clinical trials or scale manufacturing was happening then? That's a good question, worth examining at some point. cursory review of some of the investment rounds indicates wide and very international involvement of a variety investors from US, Europe, UK, Australia, South Africa, mainland China, Hong Kong and Singapore among others. These likely included many government actors: "sovereign" funds, pension funds and the like who often do these investments by allocating money to "private venture funds" (limited partners in a private venture funds are confidential). Maybe I will do a separate article on this at a later date.

Note, many people ask me "what about China and Russia?" when I talk about our own government and DoD engaged in mass genocide of Americans. I answered about China - they are allied with the US DoD on this. The CCP is profiting from the financial windfall of the US government printing dollars and throwing them into the mRNA furnaces where they are driving masses of the brainwashed citizens to suicide themselves. China claims to use "traditional vaccines" - if you believe what the Chinese say, I have a bridge to sell you.

I have not seen evidence of any similar alliance with Russia. This makes sense, because ultimately this boils down to the war of US vs Russia using proxies and alliances (as it always does). This does not mean that Russia are "the good guys". Simply that the owners of Russia (whoever they are, not necessarily based in Russia) disagree with the owners of the US (whoever they are, not necessarily based in the US). Russia is running the same "covid script", using knock off RNA/DNA injections, probably buying materials from the same suppliers, and also using war to kill off their own younger population. It's just that they are doing it for THEIR OWN interests, not that of the US and their allies.

Back to this western continent - we have already established that "Covid-19 vaccines" are biowarfare agents, legally not medicines, not pharmaceuticals, and not regulated as such.

Use of Emergency Use Authorized (EUA) covered countermeasures under a declared Public [Health](#) Emergency cannot constitute a clinical investigation (21 USC 360bbb-3(k)), therefore these countermeasures could not be tested for safety or efficacy in accordance with US law (21 CFR 312 and 21 CFR 601), nor could compliance with current Good Manufacturing Practices (cGMP) or Good Distribution Practices (GxP in general) be enforced by the FDA.

This legal fact was known to the US Government, DOD, BARDA, FDA, CDC, HHS officials signing the contracts, involved in the OWS, and it was also known to Pfizer, Moderna and other pharma companies. mRNA technology has always been designated dual-use, a category of bioweapons:

Due Diligence and Art

mRNA Injections as a Dual-Use Technology – Assessment of Threat of Misuse as Biological and Chemical Weapons.

In politics, diplomacy, and export control, "dual-use" refers to technology that can be used for both peaceful and military aims. mRNA technology, including embodiments as injectable drugs or vaccine products, has been long identified as a dual-use technology. See references...

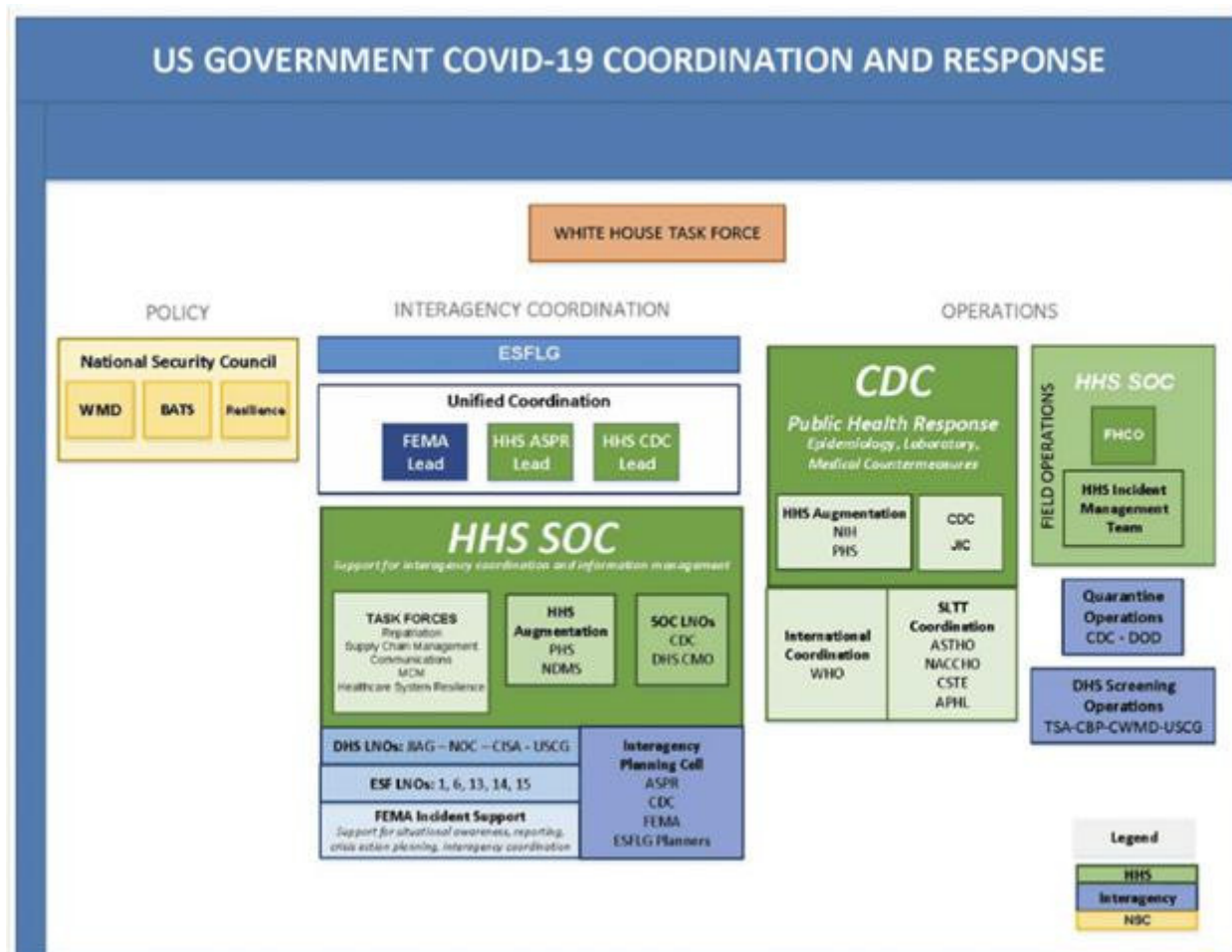
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a month ago · 59 likes · 31 comments · Sasha Latypova

From the start, "covid pandemic" was treated by the US Government as a national security matter (**i.e. war**) and covid policy was set by the National Security Council (assemblage of Defense and Intelligence heads), not HHS.

March 13, 2020: **"PanCAP Adapted U.S. Government COVID-19 Response Plan" (PanCAP-A)** states that United States policy in response to SARS-CoV-2 is set not by the public [health](#) agencies designated in pandemic preparedness protocols (Pandemic and [All Hazards Preparedness Act](#), [PPD-44](#), BIA), but rather by the National Security Council, or NSC. NSC does not have regular attendees from public [health](#) agencies and its focus is national security and foreign policy matters."

Below is the organization chart from the PanCAP-A document, p.9:



When a **known weaponizable tech** is given a liability-free, extrajudicial status shielded from all regulations, it's not hard to put 2 and 2 together. The national security, DoD and Intelligence officials absolutely knew all of this. They went ahead and authorized a \$10 billion purchase [order](#) of this weaponizable tech from the Fosun-Pfizer-BioNTech enterprise (backed

by numerous foreign governments including the Chinese), to deliver and deploy it onto Americans, **during the time of war.**

I think by now it should become clear that the “5th gen warfare” is not just the use of psyops and total control of social media by the FBI and CIA (that’s so last century!) It’s also not “profits over safety”, “bad FDA overlooked myocarditis” or “big pharma pays politicians for election campaigns”. We are way, way past that. I keep pointing out that if the motive were JUST PROFIT, then the most profitable strategy would have been to ship placebo. They would not be violating any laws by doing so, there would be no adverse events and deaths, the product would look perfectly cGMP compliant, while covid would have gone away quickly by itself. Yet, the governments (plural)-pharma cartel insists on killing and injuring millions of people, obviously limiting the profit potential by doing so.

The current war is the war of the global governments (plural), that only pretend to be at odds with each other, marketing themselves as “left”, “right”, “communists”, “green”, “capitalists”, “socialists”, “populists”, “conservatives”, etc. etc. in a never ending clown show of the political theater. Behind the scenes, the “official enemies” are partners and co-investors into “joint ventures” against us, people of the world. They use taxpayers’ money to fund, develop, then “approve”, purchase and deploy prohibited biowarfare agents for killing and injuring **their own civilian population, their own armed forces, first responders, healthcare workers, pregnant women and children.** To stop this every one of us must start using correct precise language, start calling things what they really are.

https://sashalatypova.substack.com/p/the-role-of-the-us-dod-and-their-publication_id=870364&post_id=91532704&isFreemail=true

American Domestic Bioterrorism Program

Building the case to prosecute members of Congress, presidents, HHS and DOD secretaries and federal judges for treason under 18 USC 2381.



[Katherine Watt](#)

Apr 28, 2022

Research and organizing tool first posted April 28, 2022, subject to ongoing revision as new information comes to light. Last updated Dec. 22, 2022.

Other formats: [Sept. 2022 small-print PDF](#) (67 pages); [Sept. 2022 large-print PDF](#) (101 pages); Nov. 2022 [Statutory History Orientation Memo](#) (45 pages); Dec. 2022 [Legal Structures Outline](#) (2 pages), Dec. 2022 [Six Statutory Frameworks - Executive Summary](#) (14 pages)

OVERVIEW

I started looking closely at the legal architecture supporting the Covid national prison [panopticon](#) on Jan. 30, 2022, after hearing [Attorney Todd Callender's interview](#), which provided information about the American domestic legal framework; how it fit with the oddly-coordinated pandemic story told by governments worldwide; and how it relates to the World [Health](#) Organization International [Health](#) Regulations of 2005 at the center.

I wrote up the interview:

- [Legal Walls - Short Version](#)
- [Legal Walls of the Covid-19 Kill Box](#); [PDF](#)

Prior to that day, I'd spent a lot of time, with increasing confusion and alarm and despair, trying to figure out why the U.S. Constitutional legal system hadn't put a stop to the nonsense as its nonsensicality became obvious to so many people.

Why did it continue, with no end in sight, and not even a glimpse of a path to the end?

Since then, as I've dug into Callender's analysis following the supporting paper trails, I've learned why, and how.

A whole lot of things that once were federal and state crimes and civil rights violations have been legalized by Congress through legislative, statutory revisions to the United States Code, signed by US Presidents, and implemented at the administrative, regulatory level by the Department of [Health](#) and Human Services and Department of Defense through the Code of Federal Regulations.

I've reported on those findings in small bits and pieces, connecting the laws to court cases, executive orders, guidance documents for industry and researchers, academic papers, intellectual property patents, regulatory amendments, psychological manipulation programs, geopolitical developments and other facts as they've floated across my field of view.

I think the critical decay began around 1983, when the 'public [health](#) emergencies' section was added to the 1944 Public [Health](#) Service Act, although the 1944 PHSA itself represented an additional militarization of human medicine in the United States.

Most of the worst laws have been passed since 2000 — just before 9/11 and the US Department of Defense false flag anthrax attacks.

They are listed below, with links to the full text of each law, and a short summary of what I understand about how each one fits into the overall scheme.

The basic goal of the architects, which has been achieved, was to set up legal conditions in which all governing power in the United States could be automatically transferred from the citizens and the three Constitutional branches into the two hands of the [Health](#) and Human Services Secretary, effective at the moment the HHS Secretary himself declared a public [health](#) emergency, legally transforming free citizens into enslaved subjects.

That happened on [Jan. 31, 2020, in effect as of Jan. 27, 2020](#) through the present day.

In other words: **Congress and US Presidents legalized and funded the overthrow of the U.S. Constitution, the U.S. government and the American people, through a massive domestic bioterrorism program relabeled as a public [health](#) program, conducted by the HHS Secretary and Secretary of Defense on behalf of the World [Health](#) Organization and its financial backers.**

Navigation Tool/Jump To:

- [1900-1929](#)
- [1930-1939](#)
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Related reporting:

- [COVID-19 injectable bioweapons as case study in legalized, government-operated domestic bioterrorism](#)
- [Faked Clinical Trials and 'Real World Evidence'](#)
- [US federal crimes for which there is evidence to prosecute Covid-19 bioterrorists who occupy US government positions](#)
- [22 worst Congressional bioterrorism authorization and funding laws passed since 1983](#)
- [Run-up to the American bioterrorist State's Jan. 31, 2020 declaration of war - Part 1](#) (2014-2017)

- [Run-up to the American bioterrorist State's Jan. 31, 2020 declaration of war - Part 2](#) (2018-2020)
 - [Timeline \(1819-2022\) of Supreme Court cases](#), related state cases and treaties
-

1900-1929 - Presidents Theodore Roosevelt, William Howard Taft, Woodrow Wilson, Warren Harding, Calvin Coolidge, Herbert Hoover

- 1907 - [Treaty at the Hague](#) — Convention Respecting the Laws and Customs of War on Land, including Section III, Military Authority Over the Territory of the Hostile State: “[Art. 42](#). Territory is considered occupied when it is actually placed under the authority of the hostile army.”
 - 1909 - Launch of the [Round Table Movement](#). “By 1919, the Round Table Movement changed its name to the *Royal Institute for International Affairs* (aka: Chatham [House](#)) with the Round Table name relegated to its geopolitical periodical... in America, where knowledge of the British Empire’s subversive role was more widely known, the name “American Institute for International Affairs” was still too delicate. Instead the name Council on Foreign Relations” was chosen and was chartered in 1921.”
 - 1913/12/23 - US Congress and President Wilson passed Federal Reserve Act. PL 63-43, [38 Stat. 251](#). Created Federal Reserve Bank, central banking system in United States.
 - 1921/11/23 - US Congress and President Harding passed Sheppard-Towner Maternity and Infancy Protection Act. PL 67-97, [42 Stat. 224](#). Established status of American-born babies — human beings — as collateral for national debt owed to international bankers; program operated through birth certificates/security bonds filed with state registries of vital statistics. Expired 1929, replaced by 1935 Social Security Act.
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1930-1939 - Presidents Herbert Hoover, Franklin D. Roosevelt

- 1933/04/05 - President Roosevelt signed [Executive Order 6102](#), under state of emergency (Great Depression). Ratified by Congress through [House](#) Joint Resolution 192. Forbade the hoarding 'of gold or silver coin or bullion or currency,' confiscated gold held by private individuals, to remove the constraint on the Federal Reserve (1913 Federal Reserve Act) preventing it from increasing the money supply.

- 1933/06/05 - Congress passed [House Joint Resolution 192](#), ratifying President Roosevelt's Executive [Order](#) 6102; declared bankruptcy of US government; suspended gold standard; pledged lives of American people (registered at birth through Social Security program) as collateral/debt slaves to international bankers, against national debt.
 - 1933/06/12 - London Economic Conference began. Report on Matthew Ehret, *Clash of the Two Americas: Open vs. Closed Systems Collide*: [How Roosevelt Halted Previous Attempts to Implement a New World Order](#).
 - 1935/08/14 - US Congress and President Roosevelt passed Social Security Act - PL 74-271. [49 Stat. 620](#). Social Security Act governs Medicare and Medicaid, two of the federal authorization and funding pathways through which 'breakthrough' devices and drugs, fast-track products, products eligible for accelerated approval and other FDA-classified products are developed, manufactured and used on humans. Amendments to SSA since 1983 and pending, have expanded/will further expand the novel drug and device/bioweapon classes eligible for fast-tracked federal research and deployment funding within the Medicare/Medicaid programs.
 - 1938/06/25 - Congress and President Roosevelt passed **Federal Food Drug and Cosmetic Act** (FDCA). PL 75-717, [52 Stat. 1040](#). Original stated purpose: "to prohibit the movement in interstate commerce of adulterated and misbranded food, drugs, devices, and cosmetics." Codified at 21 USC 9. By the outbreak of Covid in late 2019, FDCA had been amended by several decades of Congressional acts to become one of the key laws under which the American domestic bioterrorism program is authorized, funded and operated.
 - 1939/09/01 - Globalists launched World War II.
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1940-1949 - Presidents Franklin D. Roosevelt, Harry S. Truman

- 1944/07/01 - 07/22 - Globalists negotiated [Bretton Woods Articles of Agreement](#) to establish a centralized global financial and banking system.
- 1944/07/01 - Congress and President Roosevelt passed **Public [Health](#) Service Act** (PHSA). PL 78-410, [58 Stat. 682](#). Consolidated, centralized and militarized the American public [health](#) system that had developed within several agencies since the Revolution. Codified at 42 USC 201.
- 1945/04/12 - President Roosevelt died; President Truman took [office](#).
- 1945/07/31 - Congress and President Truman passed Bretton [Woods](#) Agreement Act, PL 79-171, [59 Stat. 512](#), authorizing President to accept membership in International

Monetary Fund and International Bank for Reconstruction and Development, later known as World Bank.

- 1945/09/02 - Globalists ended World War II.
- 1945/10/24 - Globalists established United Nations. US Congress ratified treaty.
- 1945/11/20 - Globalists began Nuremberg trials.
- 1945/12/27 - Bretton [Woods](#) Agreement entered into force.
- 1945/12/29 - Congress and President Truman passed International Organizations Immunities Act, PL 79-291, [59 Stat. 669](#). Corey Lynn report - [Laundering with Immunity: The Control Framework](#), Sept. 29, 2022.
- 1946/06/11 - Congress and President Truman passed Administrative Procedures Act, PL 79-404. [60 Stat. 237](#). Established framework for the administrative state to operate within a *de facto* executive branch dictatorship, through the “committed to agency discretion” override of both the legislative process and judicial review. Codified at 5 USC 551.
- 1946/07/22 - Globalists established the World [Health](#) Organization and adopted the WHO Constitution, signed by 61 nations at International [Health](#) Conference in [New York](#), to enter into force as of 04/07/1948. WHO Constitution amendments passed by World [Health](#) Assembly 02/03/1977 ; 01/20/1980 ; 07/11/1994 ; 09/15/2005.
- 1946/10/01 - Globalists concluded Nuremberg trials.
- 1947 National Security Act - 61 Stat. 499. Set up precursors to Federal Emergency Management Agency (FEMA).
- 1947/10/30 - Globalists adopted General Agreement on Tariffs and Trade (GATT) treaty.
- 1948 - UN Universal Declaration of Human Rights, part of International [Bill](#) on Human Rights
- 1948 US Information and Educational Exchange Act (Smith-Mundt). PL 80-402. 62 Stat. 6. Set up programs for US propaganda distribution in foreign countries; limited use of government propaganda on American population. ‘Modernized’ to authorize domestic propaganda in 01/02/2013 National Defense Authorization Act.
- 1948/01/01 - General Agreement on Tariffs and Trade (GATT) treaty entered into force.
- 1948/04/07 - World [Health](#) Organization Constitution entered into force.
- 1948/06/14 - Congress authorized President Truman to accept membership in World [Health](#) Organization on behalf of US government. PL 643, 64 Stat. 441. Codified at [22 USC 290](#).
- 1949/04/04 - US Senate ratified North Atlantic Treaty Organization (NATO) treaty.
- 1949/06/18 - George Orwell published *1984*.
- 1949/08/24 - NATO treaty entered into force.
- 1949 - Geneva Conventions

1950-1959 - Presidents Harry Truman, Dwight Eisenhower

- 1951/05/25 - Globalists adopted first International Sanitary Regulations at the World [Health](#) Organization World [Health](#) Assembly, to enter into force 10/01/1952. International Sanitary Regulations were revised and renamed International [Health](#) Regulations in 1969. Revised again 1973, 1981, 2005. Draft revisions under review 2022.
- 1951 - Globalists adopted UN Convention on the Prevention and Punishment of the Crime of Genocide.
- 1952/09/14 - Roman Catholic Pope Pius XII presented speech [On the Moral Limits of Medical Research and Treatment](#) to First International Congress on Histopathology of the Nervous System. "Insofar as the moral justification of the experiments rests on the mandate of public authority, and therefore on the subordination of the individual to the community, of the individual's welfare to the common welfare, it is based on an erroneous explanation of this principle. It must be noted that, in his personal being, man is not finally ordered to usefulness to society. On the contrary, the community exists for man."
- 1952/09/27 - President Truman signed Executive [Order](#) 10399 establishing the US Surgeon General as the "[health](#) administrator" for the World [Health](#) Organization on American soil, under 1948 WHO Constitution and 1951 WHO International Sanitary Regulations. [17 Federal Register 8648](#).
- 1952/10/01 - WHO International Sanitary Regulations of 1951 entered into force in WHO member states.
- 1953/03/12 - President Eisenhower transmitted Reorganization Plan No. 1 of 1953 to Congress, subordinating US sovereignty to WHO International Sanitary Regulations, to be implemented by Surgeon General through the Department of [Health](#), Education and Welfare (later renamed [Health](#) and Human Services). [18 Federal Register 2053](#). Codified at 42 USC 202.

1960-1969 - Presidents Dwight Eisenhower, John F. Kennedy, Lyndon Johnson, Richard Nixon

- 1961/01/17 - President Eisenhower delivered [Farewell Address](#), warning Americans of the military-industrial-Congressional complex and the "danger that public policy could itself become the captive of a scientific-technological elite."

- 1962/10/11 - Roman Catholic Pope John XIII convoked Second Vatican Council (Vatican II). Through the council, Satanic globalists expanded and deepened their infiltration to [destroy the institutional Catholic Church and weaken Catholic faith around the world](#).
- 1963/06/30 - [Enthronement of Lucifer ceremony](#) coordinated with consecration of Pope Paul VI.
- 1963/11/22 - President Kennedy assassinated; President Johnson took [office](#).
- 1964/06 - Globalists adopted the [Declaration of Helsinki](#) on ethics of human experimentation, through World Medical Association. [Revised seven times since](#): 1975, 1983, 1989, 1996, 2000, 2008, 2013.
- 1965/12/08 - Roman Catholic Pope Paul VI concluded Second Vatican Council.
- 1966/04/25 - President Johnson transmitted Reorganization Plan No. 3 of 1966 to US Congress, transferring US Surgeon General's authorities to Secretary of [Health](#), Education and Welfare department, effective 06/25/1966. [31 Federal Register 8855](#).
- 1968/04/04 - Assassination of Martin Luther King Jr.
- 1968/06/06 - Assassination of Robert F. Kennedy.
- 1968/07/25 - Roman Catholic Pope Paul VI issued papal encyclical [Humanae Vitae](#) on meaning of human life, and Catholic prohibition of abortion and contraception.
- 1969 - *Globalist* - WHO International Sanitary Regulations, in effect since 10/01/1952, revised and renamed International [Health](#) Regulations. Revised again 1973, 1981, 2005. Draft revisions under review 2022.
- 1969/06/09 - Dr. Donald MacArthur testified to [US Senate hearing on DOD appropriations](#), about development of "new infective microorganisms which could differ in certain important aspects from any known disease-causing organisms. Most important of these is that it might be refractory to the immunological and therapeutic processes upon which we depend to maintain our relative freedom from infectious disease."
- 1969/11/19 - Congress and President Nixon passed Armed Forces Appropriations Act. PL 91-121, [83 Stat. 209](#). Section 409 authorized Department of Defense to use human subjects for experiments in chemical and biological weapons, established reporting requirements (DOD reports to Congress) codified at 50 USC 1511(a) and authorized President to suspend informed consent and other provisions during a declared war or national emergency, codified at 50 USC 1515. Congressional reporting requirements amended 1977 and 1982, repealed 1996.
- 1969/11/25 - President Nixon [Statement on Chemical and Biological Defense Policies and Programs](#)
- 1969/11/30 - New [Ordo Missae](#), "liturgical innovation," introduced by Pope Paul VI, breaking the tradition of centuries.

1970-1979 - Presidents Richard Nixon, Gerald Ford, Jimmy Carter

- 1970 - Globalists, through Club of Rome, published [*The Predicament of Mankind: Quest for Structured Responses to Growing World-wide Complexities and Uncertainties, A Proposal*](#)
- 1970 - Zbigniew Brzezinski published *Between Two Ages: America's Role in the Technotronic Era*.
- 1970/03/16 - Congress and President Nixon passed An Act to Establish a Commission on Population Growth and the American Future. PL 91-213, [84 Stat. 67](#).
- 1970/08/15 - Congress and President Nixon passed Economic Stabilization Act of 1970. PL 91-379, [84 Stat. 799](#). Authorized President to stabilize prices, rents, wages, salaries, interest rates, dividends and similar transfers as part of a general program of price controls within the American domestic goods and labor markets. Used by Nixon in August 1971.
- 1970/10/26 - Congress and President Nixon passed Legislative Reorganization Act. PL 91-510, [84 Stat. 1140](#).
- 1970/11/01 - Roman Catholic Archbishop Marcel Lefebvre founded [Society of St. Pius X](#) to preserve traditional Catholic teachings in the wake of the Second Vatican Council.
- 1971 - Globalists, through Henry Kissinger and Klaus Schwab, established the World Economic Forum.
- 1971 - President Nixon launched the War on Drugs
- 1971/01 - Six banks in the European Community, under Jacob Rothschild's direction, consolidated into Inter-alpha Group of Banks.
- 1971/08/15 - President Richard Nixon directed the Treasury Secretary to suspend, with some exceptions, the convertibility of the dollar into gold or other reserve assets, ordering the gold window to be closed such that foreign governments could no longer exchange their dollars for gold, and issued Executive [Order](#) 11615 (pursuant to the [Economic Stabilization Act of 1970](#)), imposing a 90-day freeze on wages and prices in [order](#) to counter inflation.
- 1971/08 - US Department of [Health](#), Education and Welfare, National Institutes of [Health](#), National Cancer Institute published [Special Virus Program, Progress Report 8](#)
- 1971/12/23 - US Congress and President Nixon passed National Cancer Act. PL 92-216, [85 Stat. 778](#). Expanded US government bioweapons development and programs under pretext of cancer research.
- 1972 - Globalists, through Club of Rome, published [Limits to Growth](#), expanding on 1970 proposals in Predicament of Mankind.
- 1972 - Globalists, through Bulletin of the World [Health](#) Organization, published two-part series on *Virus-associated immunopathology: animal models and implications for human*

disease, [Part 1](#) and [Part 2](#), addressing potential of lab-developed viral, communicable bioweapons to cause cancers and other life-limiting autoimmune and immune dysregulation disorders.

- 1972/04/10 - Globalists opened [UN Convention on the Prohibition of the Development, Production and Stockpiling of Bacteriological \(Biological\) and Toxin Weapons and on their Destruction](#) for signing, leaving major loopholes for biological and toxic agents allegedly developed for 'protective' or 'prophylactic' purposes.
- 1972/08 - US Department of [Health](#), Education and Welfare, National Institutes of [Health](#), National Cancer Institute published [Special Virus Program, Progress Report 9](#)
- 1973 - Trilateral Commission
- 1973/01/22 - US Supreme Court issued ruling in *Roe v. Wade*, [410 US 113](#), on abortion, eroding moral status of human beings based on developmental status/age and finding a 'right' to abortion in the US Constitution.
- 1974/04/01 - Richard Gardner published essay in Foreign Affairs: [The Hard Road to World Order](#). "In short, the 'house of world order' will have to be built from the bottom up rather than from the top down. It will look like a great 'blooming, buzzing confusion,' to use William James' famous description of reality, but an end run around national sovereignty, eroding it piece by piece, will accomplish much more than the old-fashioned frontal assault."
- 1974/04/24 - Secretary of State Henry Kissinger promulgated National Security Study Memorandum 200, [Implications of Worldwide Population Growth for U.S. Security and Overseas Interests](#). NSSM 200 directed Secretary of Defense, Secretary of Agriculture, CIA Director, Deputy Secretary of State and Administrator for US Agency for International Development to study international political and economic implications of population growth and offer possible courses of action for the U.S. The resulting Kissinger Report was sent to President Nixon 12/10/1974.
- 1974 Disaster Relief Act. PL 93-288. Another statute creating precursors to FEMA.
- 1974/07/12 - US Congress and President Nixon passed National Research Service Award Act. PL 93-348, [88 Stat. 342](#). Title II set up a commission to study bioethics and protection of human subjects. Led to 1977 [Health](#), Education and Welfare report and 1979 Belmont Report.
- 1974/08/09 - President Nixon resigned; Gerald Ford took [office](#).
- 1974/11/21 - Roman Catholic Archbishop Marcel Lefebvre, founder of Society of Saint Pius X, published [1974 Declaration](#) on modernism and preservation of the Catholic faith against destructive assaults subsequent to Second Vatican Council.
- 1974/12/10 - Secretary of State Henry Kissinger's National Security Study Memorandum 200 (NSSM 200) study completed as the [Kissinger Report](#), establishing global depopulation as US geopolitical strategy.

- 1974/12/31 - US Congress and President Ford legalized private ownership of gold, reversing 1933 prohibition. PL 93-373.
- 1975/03/26 - [UN Convention on the Prohibition of the Development, Production and Stockpiling of Bacteriological \(Biological\) and Toxin Weapons and on their Destruction](#) entered into force. Codified in US law at 18 USC 175 in 1990. Both the UN convention and the US law left major loopholes for biological and chemical agents developed for 'protective' or 'prophylactic' purposes. World [Health](#) Organization, United Nations, World Economic Forum and US government drove the global bioterrorism program through those loopholes, through swine flu/H1N1, AIDS, anthrax, smallpox, MERS, SARS, SARS-CoV-2 and other communicable and injected pathogens.
- 1975/06 - Rockefeller Commission published [Report to the President on CIA Activities Within the US](#).
- 1975/11/26 - President Gerald Ford endorsed the Kissinger Report's depopulation plan through [National Security Decision Memorandum 314](#)
- 1976/01 - [Swine influenza/H1N1 outbreak started at Fort Dix](#); in April, Congress funded a vaccine development/mass vaccination through Merck; in late September injections began. Heart attacks, Guillain-Barre syndrome, deaths and other adverse effects resulted. In December, campaign suspended and never restarted.
- 1976/03/23 - UN [International Covenant on Civil and Political Rights](#) entered into force.
- 1976/04 - Senator Frank [Church](#) Commission published a [Report on the Foreign and Military Intelligence Activities of the United States](#) in April 1976. The [Church](#) report included, at Chapter 15-F, information about chemical and biological activities, and at Chapter 17, information about "Testing and Use of Chemical and Biological Agents by the Intelligence Community." It reported on Project Chatter, Project Bluebird/Artichoke, MK-ULTRA, MK-NAOMI and other programs through which the US Government conducted experiments on human subjects against their will and to their detriment.
- 1976/09/14 - Congress and President Ford passed National Emergencies Act - PL 94-412, [90 Stat. 1255](#). Codified at 50 USC 34. This is one of the key laws [cited](#) in George W. Bush's Sept. 14, 2001 Proclamation 7463, *Declaration of National Emergency by Reason of Certain Terrorist Attacks* and renewed every year since, most recently by Biden in Sept. 2021. It's also one of the laws cited in Donald Trump's March 13, 2020 Proclamation 9994, *Declaring a National Emergency Concerning the Novel Coronavirus Disease (COVID-19) Outbreak*, renewed every year since, most recently by Biden in Feb. 2022.
- 1977/01/14 - US Department of [Health](#), Education and Welfare published report on informed consent of human subjects of biomedical experiments, 45 CFR 46, *Protection of Human Subjects: Research Involving Prisoners and Notice of Report and Recommendations of the National Commission for the Protection of Human Subjects of*

Biomedical and Behavioral Research, in compliance with 1974 National Research Service Award Act. [42 Federal Register 3076](#).

- 1977/07/30 - Congress and President Carter passed Department of Defense Appropriations Authorization Act of 1978. PL 95-79, [91 Stat. 323](#). Section 808 addressed DOD use of military personnel as research subjects for biological and chemical weapons under 1969 law, codified at 50 USC 1520; required notice to be given to local officials before subjecting civilian populations to chemical and biological weapons tests; required DOD reporting to Congress. The provision on DOD reporting to Congress was amended in 1982 and repealed in 1996. Other provisions of the law were amended in 1997 to expand experimentation on military personnel, through the NDAA for FY1998 at Section 1078 and the Emergency Use Authorization provisions of the 1997 Food and Drug Administration Modernization Act at Section 402.
- 1979/04/18 - National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research published the [Belmont Report](#) on ethics of human subjects research, in compliance with 1974 National Research Service Award Act and informed by 1977 HEW report and recommendations.
- 1979/10/17 - Congress and President Carter passed Department of Education Organization Act. PL 96-88, [93 Stat. 668](#). Section 509 redesignated the US [Health](#), Education and Welfare Department as the [Health](#) and Human Services Department. From that point to the present, the Secretary of [Health](#) and Human Services has exercised authorities under the WHO Constitution and WHO International [Health](#) Regulations, as transferred from Surgeon General to HEW Secretary in 1966.

1980-1989 - Presidents Ronald Reagan, George H.W. Bush

- 1980 Comprehensive Environmental Response, Compensation and Liability Act. PL 96-510, 94 Stat. 2767. Superfund Act. Set up federal programs for cleanup of toxic chemical dumpsites.
- 1980/06/16 - US Supreme Court ruling in *Diamond v. Chakrabarty*, [447 US 303](#). Held: A live, human-made micro-organism is patentable subject matter under 35 USC 101.
- 1981/06/01 - HHS-Food and Drug Administration Final Rule *Protections for Human Subjects; Prisoners Used as Subjects in Research*, 21 CFR 50, implementing 1979 recommendations of National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research, went into effect. [45 Federal Register 36386](#)

- 1981/07/27 - HHS-FDA Final Rule *Protection of Human Subjects; Informed Consent* (21 CFR 50.20) and *Protection of Human Subjects; Standards for Institutional Review Boards for Clinical Investigations* (21 CFR 56.101) went into effect. [46 Federal Register 8942](#). Both were amended many times thereafter.
- 1982 - Roussel-Uclaf developed RU-486/mifepristone chemical abortion pill. Approved by US FDA in Sept. 2000.
- 1982/12/21 - Congress and President Reagan passed Congressional Reports Elimination Act. PL 97-375, [96 Stat. 1822](#). Section 203(a) amended requirements for DOD report to Congress on use of human subjects in chemical and biological weapons research under 50 USC 1511(a). Reporting requirement repealed by Congress, 02/10/1996, PL 104-106 at Section 1061(k).
- 1983/07/13 - Congress and President Reagan passed Public [Health](#) Service Act Amendment. PL 98-49, [97 Stat. 245](#). Section 319 amended Public [Health](#) Service Act to add a 'Public [Health](#) Emergencies' program, granting new powers to [Health](#) and Human Services Secretary and establishing a \$30 million slush fund called the Public [Health](#) Emergencies Fund. Codified at 42 USC 247d. [Summary posted April 20, 2022](#).
- 1983/12/22 - President Reagan signed Executive [Order](#) 12452, listing communicable diseases subjecting citizens to forcible apprehension and detention under [Health](#) and Human Services Secretary's quarantine authority through PHSA, [42 USC 264b](#), including "Cholera or suspected Cholera, Diphtheria, infectious Tuberculosis, Plague, suspected Smallpox, Yellow [Fever](#), and suspected Viral Hemorrhagic Fevers (Lassa, Marburg, Ebola, Congo-Crimean, and others not yet isolated or named)." [48 Federal Register 56927](#)
- 1985/11/20 - Congress and President Reagan passed [Health](#) Research Extension Act. PL 99-158, [99 Stat. 877](#). Section 498 prohibited HHS from funding or conducting fetal tissue research for three years. Codified at 42 USC 299g.
- 1986/03/29 - Robert Strecker delivered to Congress and published report on AIDS outbreak: [This Is a Bioattack Alert](#). Report connected US government cancer virus research to virus-induced immune system disorders and cancer in AIDS patients.
- 1986/07/13 - Congress and President Reagan passed Superfund Amendments and Reauthorization Act. PL 99-499, [100 Stat. 1613](#). Title III, Emergency Planning and Community Right to Know Act related to toxic chemicals and federal government authority.
- 1986/08/27 - Roman Catholic Archbishop Marcel Lefebvre published [Letter to 8 Cardinals Regarding the Assisi Affair](#), addressing dangers to the Catholic faith presented by Pope John Paul II's planned Interfaith Peace Service.
- 1986/09/18 - Roman Catholic Pope John Paul II conducted multi-religious Interfaith Peace Service in Assisi, Italy.

- 1986/11/14 - Congress and President Reagan passed State Comprehensive Mental [Health](#) Services Plan Act - PL 99-660, [100 Stat 3743](#). Title III, National Childhood Vaccine Injury Act, amended Public [Health](#) Service Act to establish and fund a National Vaccine Program; grant vaccine manufactures legal immunity for injuries and deaths caused by their products; establish and fund a tax revenue/debt-funded National Vaccine Injury Compensation Program. Codified at 42 USC 300aa.
- 1986/12/02 - Roman Catholic Archbishop Marcel Lefebvre and Bishop Antonio de Castro Mayer published [Joint Declaration Against Assisi](#), again deploring the weakening of the Catholic faith by Vatican leaders under the influence of the Second Vatican Council.
- 1987/06/27 - UN [Convention against Torture and Other Cruel, Inhuman or Degrading Treatment or Punishment](#), drafted in 1984, signed 1985, entered into force.
- 1988/11/04 - Congress and President Reagan passed Genocide Convention Implementation Act of 1987, PL 100-606, [102 Stat. 3045](#), to implement the International Convention on the Prevention and Punishment of Genocide. Codified at 18 USC 1091.
- 1988/11/04 - Congress and President Reagan passed [Health](#) Omnibus Programs Extension Act. PL 100-607, [102 Stat. 3048](#). Section 105 established National Center for Biotechnology Information under Public [Health](#) Service Act (42 USC 286c). Section 156 extended fetal tissue research moratorium imposed in 1985 for two more years. Section 201 outlined and funded HIV-AIDS research under direction of NIH/NIAID/Fauci (42 USC 300cc). Section 256 increased funding for the Public [Health](#) Emergencies Fund to \$45 million (42 USC 247d).
- 1988/11/23 - Congress and President Reagan passed Robert T. Stafford Disaster Relief and Emergency Act. PL 100-707, [100 Stat. 4689](#). Amended 1974 Disaster Relief Act, FEMA law; redefined 'emergency' and 'major disaster;' established procedures for Presidential disaster and emergency declarations, DOD domestic deployment of military and more. Codified at 42 USC 5121.
- 1989/12/19 - Congress and President George H.W. Bush passed Omnibus Budget Reconciliation Act. PL 101-239, [103 Stat. 2106](#). Section 6601 amended Vaccine Injury Compensation Program, set up special master program.

1990-1999 - Presidents George H.W. Bush, William J. Clinton

- 1990/05/22 - Congress and President Bush passed Biological Weapons Antiterrorism Act of 1989. PL 101-298, [104 Stat. 201](#). Drafted by Francis Boyle to bring US into compliance with 1975 UN convention. Establishing as [criminal](#), acts of those who "knowingly develops, produces, stockpiles, transfers, acquires, retains, or possesses any

biological agent, toxin, or delivery system for use as a weapon, or knowingly assists a foreign state or any organization to do so,” and defined ‘for use as a weapon’ to “not include the development, production, transfer, acquisition, retention, or possession of any biological agent, toxin, or delivery system for prophylactic, protective, or other peaceful purposes.” Codified at 18 USC 175.

- 1990/12/21 - HHS Interim Final Rule: *Informed Consent for Human Drugs and Biologics; Determination that Informed Consent is Not Feasible* - [55 Federal Register 52814](#)
- [1991 Common Rule](#) governing research on human subjects.
- 1992/06/03 - United Nations opened UN Conference on Environment and Development, commonly called the Earth Summit, in Rio de Janeiro, Brazil.

179 participating nations adopted [Agenda 21 \(later renamed Agenda 30\)](#), laying out plans for depopulation, elimination of private property, and elimination of borders and national sovereignty. Implicitly defined living human beings as biological weapons of mass destruction, against which lethal chemical and biological agents could be construed as ‘protective’ and ‘prophylactic’ and therefore exempt from 1975 UN Convention on Prohibition of Biological Weapons. UN Framework Convention on Climate Change opened for nation-state signatories to sign.

- 1992/07/10 - Congress and President Bush passed Alcohol, Drug Abuse, Mental [Health](#) Administration (ADAMHA) Restructuring Act. PL 102-321, [106 Stat. 323](#). Expanded drug abuse prevention and treatment programs; reorganized HHS subdivisions.
- 1992/10/27 - Congress and President Bush passed Preventative [Health](#) Amendments. PL 102-531, [106 Stat. 3504](#). Changed name from Centers for Disease Control to Centers for Disease Control and Prevention.
- 1993/06/10 - Congress and President Clinton passed National Institutes of [Health](#) Revitalization Act, PL 103-43, [107 Stat. 122](#). Reorganized and expanded research programs; reversed moratorium on fetal tissue research.
- 1993/11/16 - Congress and President Clinton passed Religious Freedom Restoration Act. PL 103-141, [107 Stat. 1488](#). Affirmed Constitutional protections for free exercise of religion under First Amendment. Related to military personnel requests for religious exemptions from vaccine mandates, not accepted by DOD. Codified at 42 USC 2000bb.
- 1993/11/30 - Congress and President Clinton passed NDAA for FY1994, PL 103-160, [107 Stat. 1547](#). Section 1703 related to DOD reporting to Congress on chemical and biological weapons testing programs. Codified at 50 USC 1523. Amended 11/18/1997 and 10/17/2006. Repealed 12/23/2016, effective 12/31/2021?? , Also authorized DOD to “enter into agreements with Secretary of HHS to provide support for vaccination programs...in the US through use of the excess peacetime biological weapons defense capability of the DOD.” Codified at 50 USC 1524.

- 1994/03/21 - United Nations [Framework Convention on Climate Change](#) entered into force.
- 1994/09/05 - United Nations opened the International Conference on Population and Development in Cairo, Egypt. 179 nation-states signed on to a 20-year Programme of Action for depopulation, which was [extended in 2010 to cover 2014-2034](#).
- 1994/09/13 - Congress and President Clinton passed Violent Crime Control and Law Enforcement Act (Clinton Crime [Bill](#)). PL 103-322, [108 Stat. 1796](#). Expanded American prison state, by expanding predicates for incarcerating nonviolent civilians for long sentences, increasing funding for prison construction/operation, and law enforcement officers.
- 1994/12/08 - Rockefeller Senate Report on US government chemical and biological weapons research, development, testing and deployment programs. [S.Prt. 103-97](#).
- 1995 - Launch of World Trade Organization, update to 1947 General Agreement on Trade and Tariffs.
- 1996/02/10 - Congress and President Clinton passed National Defense Authorization Act for FY96. PL 104-106, [110 Stat. 443](#). Section 1061(k) repealed 50 USC 1511 as adopted in 1977 and amended in 1982, eliminating requirement that DOD report to Congress on chemical and biological weapons experiments conducted on military personnel.
- 1996/04/24 - Congress and President Clinton passed Antiterrorism and Effective Death Penalty Act; Illegal Immigration Reform and Immigrant Responsibility Act; Prison Litigation Reform Act. PL 104-132. [110 Stat. 1214](#). Section 521(a) prohibited DOD chemical and biological weapons testing in urban and suburban areas, codified at 18 USC 2332C. That provision was repealed in 1998. Also related to court stripping: Congress passing laws to remove federal courts' oversight power regarding legislative and executive acts, eliminate checks and balances. See ACLU report, Oct. 2001, [Upsetting Checks and Balances: Congressional Hostility Toward the Courts in Times of Crisis](#).
- 1996/12/17 - UN Comprehensive Convention on International Terrorism opened for negotiation by [resolution 51/210 forming ad hoc committee](#); subsequently deadlocked over definition of terrorism.
- 1997/04/29 - UN [Convention on the Prohibition of the Development, Production, Stockpiling and Use of Chemical Weapons and on their Destruction](#) entered into force, after drafting in 1992 and signing in 1993.
- 1997/11/18 - Congress and President Clinton passed National Defense Authorization Act for FY98 - PL 105-85, [111 Stat. 1915](#). Section 1078, "Restrictions on the use of human subjects for testing of chemical or biological agents," repealed and replaced a 1977 section of 50 USC Chapter 32, the Chemical and Biological Warfare Program. The 1977 provision (50 USC 1520) had added a requirement that DOD report to Congress about DOD human experimentation programs. In 1997, Congress replaced 1520 with 1520a,

purportedly to prohibit DOD conducting experiments on soldiers without the individual soldiers informed consent. It was passed by Congress in response to public outrage over injuries and deaths caused by mandated anthrax injections of soldiers during and after the 1991 Gulf War. However, the authority for federal government experimentation on non-consenting human beings continued; Congress simply transferred the program to the Food Drug and Cosmetics Act, 21 USC 360bbb (see below, passed three days after the NDAA) under declared emergency situations (Emergency Use Authorizations/EUA).

- 1997/11/21 - Congress and President Clinton passed Food and Drug Administration Modernization Act - PL 105-115, [111 Stat. 2296](#). Added new section to Federal Food Drug and Cosmetics Act to expand access to investigational drugs and devices during emergency situations. Codified at 21 USC 360bbb - "Expanded Access to Unapproved Therapies and Diagnostics". This was the beginning of the Emergency Use Authorization/EUA framework that culminated in the American government's psychological, social and economic coercion program aimed at universal injection of all American citizens with products marketed as Covid-19 vaccines, operational from mid-2020 to the present.
- 1998/03 - [Washington DC tabletop exercise](#) on smallpox epidemic. Used for political cover six months later to establish Strategic National Stockpile of US-government-controlled chemical and biological weapons, disguised as 'vaccines' and other 'pharmaceutical' products.
- 1998/10/17 - Congress and President Clinton passed National Defense Authorization Act for FY1999. PL 105-261, 112 Stat. 1920. Section 1401.
- 1998/10/21 - Congress and President Clinton passed Omnibus Consolidated and Emergency Supplemental Appropriations for FY1999 - PL 105-277, [112 Stat. 2681-358](#). Title II established the National Pharmaceutical Stockpile, later renamed the Strategic National Stockpile. Appropriated \$51,000,000, "to remain available until expended...for pharmaceutical and vaccine stockpiling activities at the Centers for Disease Control and Prevention." Division I, Chemical Weapons Convention Implementation Act of 1998, established prohibitions on chemical weapons. Codified at [18 USC 229](#) and [22 USC 6701](#).
- 1999/09/17 - [Death of Jesse Gelsinger](#) from early gene therapy trial.
- 1999/09/30 - President Clinton signed Executive [Order](#) 13139: *Improving [Health](#) Protection of Military Personnel Participating in Particular Military Operations*. Authorized administration of experimental, FDA-unapproved vaccines to members of the armed forces without informed consent. [64 Federal Register 54175](#)
- 1999/10/05 - HHS Interim Final Rule - *Human Drugs and Biologics; Determination That Informed Consent Is NOT Feasible or Is Contrary to the Best Interests of Recipients;*

Revocation of 1990 Interim Final Rule; Establishment of New Interim Final Rule. [64 Federal Register 54180](#)

- 1999/11 - Population-control zealot [Bill Gates launched GAVI](#) (Global Alliance for Vaccines and Immunizations) with \$750 million investment from [Bill](#) & Melinda Gates Foundation. Public-private partnership organization develops, tests, manufactures and deploys pharmaceutical products in low and middle-income countries.
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2000 - 2009 - Presidents William Clinton, George W. Bush, Barack H. Obama

- 2000/09 - FDA approved RU-486, mifepristone pill for use to terminate pregnancies: chemical abortion drug.
- 2000/09 - Project for the New American Century published [Rebuilding America's Defenses](#) report. "Advanced forms of biological warfare that can 'target' specific genotypes may transform biological warfare from the realm of terror to a politically useful tool."
- 2000/11/13 - Congress and President Clinton passed Public [Health](#) Improvement Act - PL 106-505, [114 Stat. 2314](#). Title I, Public [Health](#) Threats and Emergencies Act, reworked and expanded Section 319 of Public [Health](#) Service Act, 42 USC 247d (the Public [Health](#) Emergencies section first added in 1983). Appropriated funding and established a working group on bioterrorism 'countermeasures' research and development.
- 2001/09/11 - Terrorist airplane attacks on World Trade Center and Pentagon.
- 2001/09/14 - George W. Bush signed Proclamation 7463, *Declaration of National Emergency by Reason of Certain Terrorist Attacks*, under 1975 National Emergencies Act. Renewed every year since, most recently by Biden in Sept. 2021. [66 Federal Register 48199](#)
- 2001/09/18 - 2001/10/09 - Anthrax attacks on US Congress and media organizations.
- 2001/09/18 - Congress and President Bush passed Authorization for Use of Military Force. PL 107-40; [115 Stat. 224](#). Passed under the 1973 War Powers Act, 50 U.S. Code § 1541, and construed as putting the United States in a permanent state of war (Global War on Terror) with no limitations in time or geographically.
- 2001/09/23 - President Bush signed [Executive Order 13224](#), blocking property ownership and prohibiting transactions with persons who commit, threaten to commit or support terrorism. List maintained by [Office](#) of Foreign Assets Control, US Dept. of Treasury.
- 2001/10/23 - [Model State Emergency Health Powers Act](#) promulgated by CDC and the Center for Law and the Public's [Health](#) at Georgetown and Johns Hopkins Universities,

“structured to reflect 5 basic public [health](#) functions to be facilitated by law: (1) preparedness, comprehensive planning for a public [health](#) emergency; (2) surveillance, measures to detect and track public [health](#) emergencies; (3) management of property, ensuring adequate availability of vaccines, pharmaceuticals, and hospitals, as well as providing power to abate hazards to the public's [health](#); (4) protection of persons, powers to compel vaccination, testing, treatment, isolation, and quarantine when clearly necessary; and (5) communication, providing clear and authoritative information to the public.”

- 2001/10/26 - Congress and President Bush passed Uniting and Strengthening America by Providing Appropriate Tools Required to Intercept and Obstruct Terrorism (USA PATRIOT) Act - PL 107-56, [115 Stat. 272](#). Amended 18 USC 2331 - Definitions section of 18 USC 113B - Terrorism - to add “domestic terrorism,” defined as activities that “(A) involve acts dangerous to human life that are a violation of the [criminal](#) laws of the United States or of any State; (B) appear to be intended—(i) to intimidate or coerce a civilian population; (ii) to influence the policy of a government by intimidation or coercion; or (iii) to affect the conduct of a government by mass destruction, assassination, or kidnapping; and (C) occur primarily within the territorial jurisdiction of the United States.” There is [plenty of evidence to prosecute and convict Fauci, Baric, Gates, Daszak and others under this criminal statute](#). However, this is also why the conspirators used the FBI to infiltrate the January 6, 2021 [Washington](#) DC election protests, to ensure breach of the Capitol and subsequent arrests and indefinite detentions of non-violent trespassers, to create predicates to [steer and shape national panic about domestic terrorism exclusively defined as civilians challenging the legitimacy of government officials and acts](#), to steer public anger and distrust away from government agents killing, maiming and imprisoning civilians.
- 2002/06/12 - Congress and President Bush passed Public [Health](#) Security and Bioterrorism Preparedness and Response Act - PL 107-188, [116 Stat. 594](#). Major amendments to Public [Health](#) Service Act (42 USC 201) and Federal Food Drug and Cosmetics Act (21 USC 9). This law fully constructed and expanded funding for the federal government's domestic bioterrorism apparatus headquartered at the CDC, disguising it as a program to protect Americans from non-state actors. Sections included National Preparedness and Response Planning, Coordinating, and Reporting; Strategic National Stockpile; Development of Priority Countermeasures (i.e. fast-tracking approval of drugs and devices without standard safety testing, efficacy testing, and regulatory compliance); Improving State, Local, and [Hospital](#) Preparedness for and Response to Bioterrorism and Other Public [Health](#) Emergencies; Emergency Authorities (i.e. federal quarantine power); Controls on Dangerous Biological Agents and Toxins (Title II, Subtitle B: [Agricultural Bioterrorism Protection Act of 2002](#)); Safety and Security of Food and Drug Supply; Drinking [Water](#) Security and Safety. Coincidentally also in 2002, HHS-NIH-

funded (grant no. AI23946-08) [University](#) of North Carolina researcher and Fauci colleague Ralph Baric filed a [US patent \(7,279,372\)](#) on methods to make bat coronaviruses more lethal to humans, noting that “the US government has certain rights to this invention.” [More on that.](#)

- 2002/11/25 - Congress and President Bush passed Homeland Security Act - PL 107-296, [116 Stat. 2135](#). Established Department of Homeland Security as a cabinet-level administrative arm of the executive branch. Expanded militarization of domestic surveillance and law enforcement. Title V: established a Directorate of Emergency Preparedness and Response within Department of Homeland Security, headed by an Undersecretary. Strengthened crosslinks between DHS and other federal agencies: [Health](#) and Human Services, Federal Emergency Management Agency (FEMA), Department of Defense, Department of Justice and Department of Agriculture, to build and operate a public-[health](#)-predicated martial law system.
- 2003/04/04 - Congressional hearing held on [Project Bioshield: Contracting for the Health and Security of the American Public](#). Congress members discussed authorizing HHS to waive informed consent during declared emergencies. ([06/14/2022 Bailiwick post](#) with partial transcript.)
- 2003/04/04 - President Bush signed Executive [Order](#) 13295 added symptomatic SARS to list of quarantinable communicable diseases, authorizing HHS to [order](#) apprehension and indefinite detention of Americans for contracting common respiratory illnesses under [42 USC 264\(b\)](#) and [42 CFR 70.6](#). [68 Federal Register 17255](#).
- 2003/09/16 - [Model State Public Health Act](#) published by Johns Hopkins, Georgetown and CDC, working through Turning Point Initiative/Turning Point National Collaborative. Slightly less aggressive form of Model State Emergency [Health](#) Powers Act circulated in October 2001.
- 2003/11/24 - Congress and President Bush passed National Defense Authorization Act for FY2004. PL 108-136, [117 Stat. 1392](#). Section 1603(a), created 21 USC 360bbb-3 - “Section 564 - Authorization for Medical Products for Use in Emergencies” under the EUA part of the Federal Food Drug and Cosmetics Act as amended in 1997 to add 21 USC 360bbb “Expanded Access to Unapproved Diagnostics and Therapies.” At Section 1603(b)(1), Congress added Section 1107a to the military code after 10 USC 1107, authorizing the US President to waive informed consent rights of military personnel during declared emergencies and redefining the meaning of the right to be “informed of an option to accept or refuse administration of a product.”
- 2003/12/22 - US federal court in [Doe v. Rumsfeld, 297 F Supp. 2d 119](#) (DDC 2003) addressed informed consent (10 USC 1107) and Presidential waivers (10 USC 1107a) in the anthrax vaccination campaign context. Federal court enjoined DOD from overriding service members informed consent requirements with the experimental Anthrax vaccine. Eight days later, FDA fully approved the Anthrax vaccine. That FDA decision was

vacated by the Court 10/27/2004 in *Rumsfeld II*, [341 F. Supp. 2d 1 \(D.D.C. 2004\)](#). The injunction was expanded to cover the vaccine after being granted EUA status in *Rumsfeld III*. [2005 WL 774857 \(D.D.C. April 6, 2005\)](#).

- 2004/07/21 - Congress and President Bush passed Project Bioshield Act. PL 108-276, [118 Stat. 835](#). Amendments to Public [Health](#) Service Act and Federal Food Drug and Cosmetics Act. Nullified informed consent principles under US law. Amended and expanded 21 USC 360bbb on authorization for investigational drugs and devices to be used in emergencies (Emergency Use Authorization). Established program for 'qualified countermeasure' research, procurement, contracting, manufacture, use and liability exemptions. Expanded authority of NIAID Director (Fauci). Appropriated \$640,000,000 for the Strategic National Stockpile for FY2002, \$590,000,000 for smallpox vaccine development for FY2002, and \$5,593,000,000 for "procurement of security countermeasures." Expanded HHS power to subject citizens to involuntary relocation and indefinite detention on communicable disease predicates. Expanded coordination among Secretary of [Health](#) and Human Services, Secretary of Defense and Secretary of Homeland Security.
- 2005/04/01 - President Bush signed Executive [Order](#) 13375, adding symptomatic influenza to list of quarantinable communicable diseases, authorizing HHS Secretary to use force to apprehend and detain people under [42 USC 264\(b\)](#) and [42 CFR 70.6](#).. [64 Federal Register 17299](#).
- 2005/04/02 - Death of Roman Catholic Pope John Paul II. After conclave, Pope Benedict XVI took the papacy 04/19/2005.
- 2005/07/05 - HHS FDA Draft Guidance Re: Emergency Use Authorization of Medical Products. [70 FR 38689](#).
- 2005/09/15 - World [Health](#) Assembly adopted [World Health Organization International Health Regulations 2005](#) revisions. Entered into force 06/15/2007.
- 2005/12/30 - Congress and President Bush passed Department of Defense, Emergency Supplemental Appropriations to Address Hurricanes in the Gulf of Mexico, and Pandemic Influenza Act - PL 109-148, [119 Stat. 2818](#), Division C at last 14 pages: **Public Readiness and Emergency Preparedness (PREP) Act**. Amended Public [Health](#) Service Act. Established power of Secretary of [Health](#) and Human Services, during self-declared public [health](#) emergency under Section 319, to unilaterally issue declarations recommending "manufacture, testing, development, distribution, administration, or use of one or more covered countermeasures." Codified at 42 USC 247d-6d(b). Added more detail on liability shields for pandemic and epidemic products and security countermeasures. Set pre-[suit](#) hurdle requiring HHS to first bring claims against defendants, and bar private claims until after HHS claims resolved, if and only if defendant found liable. Set liability standard at willful misconduct, "establishing a standard...more stringent than negligence in any form or recklessness," requiring proof

defendant 1) intentionally engaged in misconduct 2) proximate to victim's injury or death. Established just-following-orders defense for vaccinators and others in the chain of distribution. Established court-alternative, tax-and-debt-funded Covered Countermeasure Process Fund, similar to Vaccine Injury Compensation Fund established in 1986 for products on childhood vaccine schedule. Another provision of the DOD Supplemental Emergency Appropriation funded the Public [Health](#) and Social Service Emergency Fund (PHSSEF), a slush fund under the control of the Secretary of [Health](#) and Human Services, with \$3.3 billion to start.

- 2006/06/07 - HHS-FDA Interim Final Rule, *Medical Devices; Exception From General Requirements for Informed Consent*. [71 Federal Register 32827](#)
- 2006/09 - Department of Justice published report: [Role of Law Enforcement in Public Health Emergencies: Special Considerations for an All-Hazards Approach](#). "Depending on the threat, law enforcement's role may include enforcing public [health](#) orders (e.g., quarantines or travel restrictions), securing the perimeter of contaminated areas, securing [health](#) care facilities, controlling crowds, investigating scenes of suspected biological terrorism, and protecting national stockpiles of vaccines or other medicines."
- 2006/10/17 - Congress and President Bush passed NDAA/John Warner Defense Authorization Act for FY2007 - PL 109-364, [120 Stat. 2095](#). Section 1076 amended 1807 Insurrection Act, (10 USC 333, renumbered as 10 USC 253), providing exemptions to 1878 Posse Comitatus Act, to expand the authority of federal government to deploy US military on American soil against American citizens during "natural disaster, epidemic, or other serious public [health](#) emergency, terrorist attack or incident, or other condition in any State or possession of the United States." Repealed in NDAA for FY2008. Passed again in NDAA for FY2012.
- 2006/11/28 - HHS FDA Guidance: [Gene Therapy Clinical Trials - Observing Subjects for Delayed Adverse Effects](#)
- 2006/12/19 - Congress and President Bush passed Pandemic and All-Hazards Preparedness Act. PL 109-417, [120 Stat. 2878](#). Fulfilled many of the requirements of the [World Health Organization International Health Regulations of 2005](#), by further consolidating and centralizing power in federal [Health](#) and Human Services Secretary's hands. Created new HHS department, led by new Assistant Secretary for Preparedness and Response (counterpart to the DHS Director of Emergency Preparedness and Response position created in 2002). Established rules for coordination among HHS, Secretary of Defense, Secretary of Veterans Affairs, Secretary of Transportation and "any other relevant federal agency." Established national framework subordinating state, county, tribal and local public [health](#) and law enforcement systems to federal agencies. Expanded surveillance programs. Clarified definitions of qualified countermeasure,

security countermeasure, and infectious disease for purposes of 2004 Project Bioshield Act. Established Biomedical Advanced Research and Development Authority (BARDA) division under HHS, “to facilitate a broad-based approach to emergency medical countermeasure-related activities,” including \$1,070,000,000 appropriation. Tools included HHS authority to limit competition among manufacturers of pandemic products as defined under 2004 Project Bioshield Act.

- 2007/01/15 - Congress and President Bush passed National Institute of [Health](#) Reform Act - PL 109-482, [120 Stat. 3675](#). Reorganization, consolidation of power and funding.
- 2007/05/04 - President Bush issued [National Security Presidential Directive 51](#). US Government Continuity of Operations policy.
- 2007/06/15 - [World Health Organization International Health Regulations, 2005 Amendments](#), entered into force.
- 2007/07/01 - HHS FDA [Guidance - Emergency Use Authorization of Medical Products](#). [71 FR 41083](#). Finalized draft guidance published in Federal Register 07/05/2005.
- 2007/07/07 - Roman Catholic Pope Benedict XVI issued *Summorum Pontificum*, affirming the right of Catholic priests and faithful to celebrate the pre-1962, Traditional Latin Mass.
- 2007/09/27 - Congress and President Bush passed Food and Drug Administration Amendments Act of 2007. PL 110-85, [121 Stat. 823](#). Expanded FDA power over new product authorizations and post-marketing surveillance.
- 2007/12/28 - HHS Interim Final Rule - FDA Exceptions or Alternatives to Labeling Requirements for Products Held by the Strategic National Stockpile. Effective same day. [72 FR 73589](#).
- 2008/01/28 - Congress and President Bush passed National Defense Authorization Act for FY2008. PL 110-181, [122 Stat. 325](#). Section 1068 repealed 2007 amendments to Insurrection Act which had expanded exemptions to 1878 Posse Comitatus Act limits on US Presidents’ power to deploy the military domestically. Amendments passed again in NDAA for FY2012, again giving President power to deploy military domestically.
- 2008/07 - DOJ-CDC published [A Framework for Improving Cross-Sector Coordination for Emergency Preparedness and Response](#). Merging public [health](#) and law enforcement.
- 2009 H1N1 outbreak, first mass vaccination campaign since 1976 swine flu outbreak.
- 2009/11/18 - HHS FDA Workshop Summary: [Medical Countermeasures Dispensing: Emergency Use Authorization and the Postal Model](#). “At the workshop, participants noted that EUA has a broader use beyond enabling the use of an unapproved product or extending the use of an approved product to populations for which it was not approved. In particular, it can also be used to address labeling requirements and other challenges that arise because of constraints inherent in a public [health](#) response. **‘From a legal perspective, there are a lot of situations where EUA helps get past all those**

requirements,’ said [Susan E. Sherman, J.D., M.S., is a senior attorney with the [Office](#) of the General Counsel, HHS] ‘You can change the labeling. You can change the information. You can change the dosage. You can give it to populations for which wasn’t approved.’ ”

2010-2019 - Presidents Barack H. Obama, Donald J. Trump

- 2010/03/23 - Congress and President Obama passed Patient Protection and Affordable Care Act (ObamaCare). PL 111-148, [124 Stat. 119](#). Title VII, Biologics Price Competition and Innovation Act of 2009, related to the legal, approval/authorization, labelling and marketing differences among ‘biosimilars,’ BLA (Biologics License Application) products, and EUA products.
- 2010/07/02 - President Obama signed Executive [Order](#) 13546, *Optimizing the Security of Biological Select Agents and Toxins in the United States*. [75 Federal Register 39439](#).
- 2011/01 - HHS [FDA Guidance for Industry: Potency Tests for Cellular and Gene Therapy Products](#)
- 2011/06/24 - HHS-FDA Final Rule: *Medical Devices; Exception From General Requirements for Informed Consent*. [76 Federal Register 36989](#).
- 2011/09/16 - Congress and President Obama passed Leahy Smith America Invents Act. PL 112-29, [125 Stat. 340](#). Section 33 limited the authority of the US patent [office](#) under 35 USC 101, by prohibiting issuing of patents “directed to or encompassing a human organism.” Related to 1980 *Chakrabarty* and 2013 *Myriad* Supreme Court precedents authorizing patents on genetically-modified living organisms and modified genetic material, and government-ordered mRNA and DNA spike protein Covid injections that reverse-transcribed genetic material into human genome of recipients.
- 2011/12/31 - Congress and President Obama passed National Defense Authorization Act for FY2012 - PL 112-81, [125 Stat. 1298](#). Section 1021 codified authority for US President to [order](#) military arrest and indefinite detention of American civilians without charge or trial under 10 USC 801 et seq. (Uniform Code of Military Justice), to the extent the [2001 Authorization for Use of Military Force](#), passed under the 1973 War Powers Act, (50 U.S. Code § 1541) is construed as putting the United States in a permanent state of war (Global War on Terror) and the national emergency first declared by President Bush in 2001 is extended. It has been extended, every year since.
- 2012/07/09 - Congress and President Obama passed Food and Drug Administration Safety and Innovation Act. PL 112-144, [126 Stat. 993](#). Amendments to Federal Food,

Drug, and Cosmetic Act regarding user-fee programs for prescription drugs and medical devices, generic drugs and biosimilars, and for other purposes.

- 2013/01/02 - Congress and President Obama passed National Defense Authorization Act for FY2013. PL 112-239, [126 Stat. 1957](#). Section 1078 “modernized” Smith-Mundt Act of 1948 to authorize domestic deployment of propaganda by the US government, on the American population. Propaganda used with tremendous effect on US population to instill fear and promote behavioral compliance with government orders.
- 2013/01/29 - Congress and President Obama passed Disaster Relief Appropriations Act. PL 113-2, [127 Stat. 4](#). Division B, Sandy Recovery Act: most major FEMA overhaul since 1988 Robert T. Stafford Act.
- 2013/02/28 - Roman Catholic Pope Benedict XVI resigned. After conclave, the papacy of Pope Francis began 03/13/2013.
- 2013/03/13 - Congress and President Obama passed Pandemic and All-Hazards Preparedness Reauthorization Act. PL 113-5, [127 Stat. 161](#). Renewed and updated 2006 Pandemic and All-Hazards Preparedness Act, with amendments to Public [Health](#) Service Act and Federal Food Drug and Cosmetics Act. Added sections 564A and 564B to the FDCA to further authorize emergency use of approved products in emergencies and products held for emergency use. Amended definitions of covered countermeasures and qualified pandemic and epidemic products in Section 319F-3 of PHSA (2005 PREP Act provisions). Extended definitions to include products or technologies intended to enhance the use or effect of a drug, biological product, or device used against the pandemic or epidemic or against adverse events from these products.
- 2013/06/13 - US Supreme Court ruled on *Association for Molecular Pathology v. Myriad Genetics*, [539 US 576](#), in favor of the biotech corporation and the federal government, finding that naturally-occurring DNA is not patentable, but synthetic cDNA is patentable, under 35 USC 101. Implicates mRNA/DNA injections administered on global population starting in December 2020, reverse-transcription into human genome, and whether injected humans are chattel property of Covid-19 injection patent-holders within US government/DOD, Pfizer, Moderna, AstraZeneca and Janssen.
- 2014/07/31 - President Obama signed Executive [Order](#) 13674, adding asymptomatic, suspected SARS to list of quarantinable communicable diseases under [42 USC 264\(b\)](#) and [42 CFR 70.6](#). [79 Federal Register 75461](#)
- 2014/08/19 - HHS FDA Guidance: [Decisions for Investigational Device Exemption Clinical Investigations](#). Related to federal government’s position on legal status and regulatory control differences between Emergency Use Authorization (EUA) products, Investigational New Drugs (IND) and Investigational Device Exemptions (IDE).
- 2015/04/16 - Congress and President Obama passed Medicare Access and CHIP Reauthorization (MACRA) Act. PL 114-10, [129 Stat. 87](#). Largest changes to [health](#) care system since 2010 ObamaCare. Section 511 directed HHS to clarify how changes to

human subjects protections under 1991 Common Rule would apply to Medicare and Medicaid “clinical data registries.” Related to ‘real world evidence’ with no legal protections for human subjects, replacing traditional clinical trial procedures that did have legal protections for human subjects. Codified at...

- 2015/06 - HHS FDA Guidance: [Considerations for the Design of Early-Phase Clinical Trials of Cellular and Gene Therapy Products](#)
- 2015/08 - HHS FDA Guidance: [Design and Analysis of Shedding Studies for Virus or Bacteria-Based Gene Therapy and Oncolytic Products](#)
- 2015/11/25 - Congress and President Obama passed National Defense Authorization Act for FY-2016. PL 114-92, [129 Stat. 893](#). Section 815 added ‘prototype’ procurement contracting language (Other Transactional Authority - OTA), authorizing Department of Defense to contract with pharmaceutical corporations to produce bioweapons labeled as medical countermeasures or security countermeasures. Used to contract for production of ‘Covid-19 vaccine’ bioweapons in 2020, through Medical CBRN [Chemical Biological Radiological Nuclear] Defense Consortium program members. Codified at 10 USC 2371b, renumbered 10 USC 4022 effective 01/01/2021. First two posts on this topic: [05/25/2022](#) and [05/26/2022](#).
- 2016/09/21 - HHS Final Rule - HHS Clinical Trials Registration and Results. [81 Federal Register 64981](#)
- 2016/10/17 - Congress and President Obama passed National Defense Authorization Act for FY2017. PL 114-328, [130 Stat. 2000](#). 10 USC 111 note at 130 Stat. 2400
- 2016/10/24 - HHS Workshop Summary - [The Nation's Medical Countermeasure Stockpile: Opportunities to Improve the Efficiency, Effectiveness, and Sustainability of the CDC Strategic National Stockpile](#).
- 2016/11/04 - President Obama signed [Executive Order 13747](#): *Advancing the Global [Health](#) Security Agenda to Achieve a World Safe and Secure from Infectious Disease Threats*
- 2016/12/13 - Congress and President Obama passed 21st Century Cures Act (Cures Act 1.0) - PL 114-255, [130 Stat. 1033](#). Updated and expanded Public [Health](#) Service Act “to accelerate the discovery, development, and delivery of 21st century cures.” Section 3022 authorized ‘real world evidence’ instead of clinical trials as grounds for FDA authorizing general use of experimental products, transforming Americans into human subjects and our communities into unmonitored, unregulated experimental test sites. Sections 3023 and 3024 granted broad authority for HHS Secretary to waive or alter human subject protections and informed consent requirements, by transferring each individual human subject’s risk-benefit assessment authority to the HHS Secretary, who can preemptively decide, for all subjects collectively, without knowledge of individual [health](#) conditions or conscientious beliefs, and without the subjects’ knowledge or consent, that risk is ‘minimal.’ Codified at...

- 2016/12/23 - Congress and President Obama passed National Defense Authorization Act for FY2017. PL 114-328, [130 Stat. 2509](#). Established DOD Defense Security Cooperation Agency (DSCA) and Director of DSCA, with authority to coordinate and synchronize US military with foreign military forces, and conduct domestic military campaigns in violation of the 1878 Posse Comitatus Act. Codified at [10 USC 382](#).
- 2017/01/13 - HHS [FDA Guidance: Emergency Use Authorization of Medical Products and Related Authorities](#) (Update/revision to 07/01/2007 version). Related to federal government's position on legal status and regulatory control differences between Emergency Use Authorization (EUA) products, Investigational New Drugs (IND) and Investigational Device Exemptions (IDE).
- 2017/01/19 - HHS Final Rule - [Federal Policy for the Protection of Human Subjects](#). 82 FR 7149. Joint rule by 16 federal agencies, subsequently adopted by other agencies. Revised [1991 Common Rule](#), which had been developed based on [1947 Nuremberg Code](#) and [1978 Belmont Report](#).
- 2017/01/19 HHS Final Rule - [Control of Communicable Diseases Final Rule](#). 82 FR 6890. Set up regulations governing apprehension and detention of American people on public [health](#) quarantine pretexts.
- 2017/01/23 - Department of Homeland Security published [Biological Incident Annex to the Response and Recovery Federal Interagency Operational Plans](#). At p. 70, stated that 10 USC 382 "permits Department of Defense to provide support to the Department of Justice under certain circumstances in emergency situations involving Weapons of Mass Destruction, including biological weapons and materials."
- 2017/07/25 - HHS [FDA Guidance: IRB Waiver or Alteration of Informed Consent for Clinical Investigations Involving No More Than Minimal Risk to Human Subjects](#)
- 2017/08 - HHS [FDA Guidance: Use of Real-World Evidence to Support Regulatory Decision-Making for Medical Devices](#)
- 2017/08/18 - Congress and President Trump passed FDA Reauthorization Act - PL 115-52. [131 Stat. 1005](#)
- 2017/10 - Johns Hopkins [University](#) Center for [Health](#) Security exercise and report, [SPARS Pandemic, 2025-2028, A Futuristic Scenario for Public Health Risk Communicators](#). Chapter 13 covered how government and corporate PR representatives should handle "anti-vaccine" messages. Chapter 17 covered how they should manage public awareness and anger about vaccine injury.
- 2017/12/12 - Congress and President Trump passed National Defense Authorization Act FY 2018 - PL 115-91, [131 Stat. 1283](#). Section 716 added subsection (d) to 10 USC 1107a, re: EUA product use in military. *But* see FDCA amendment, PL 115-92 (below) passed same day, which immediately repealed 10 USC 1107a(d) while adding new FDCA section on military use of EUAs.

- 2017/12/12 - Congress and President Trump passed Act to amend FDCA EUA statute, 21 USC 360bbb-3. PL 115-92, [131 Stat. 2023](#). Provided for “Additional Emergency Uses for Medical Products to Reduce Deaths and Severity of Injuries Caused by Agents of War.” Codified at...
- 2018/01 - FEMA published [Pandemic Crisis Action Plan/PanCAP](#).
- 2018/06/19 - HHS Final Rule - Federal Policy for the Protection of Human Subjects: Six Month Delay of the General Compliance Date of Revisions While Allowing the Use of Three Burden-Reducing Provisions During the Delay Period. [83 Federal Register 28497](#)
- 2018/06/19 - [Biodefense in the Age of Synthetic Biology](#) published by US National Academies of Sciences, Engineering, Medicine.
- 2018/10/05 - Congress and President Trump passed Federal Aviation Administration Reauthorization Act. PL 115-254, [132 Stat. 3186](#). Division D, Disaster Recovery Reform Act, another major FEMA update.
- 2018/10/09 - Johns Hopkins [University](#) Center for [Health](#) Security published report [Technologies to Address Global Catastrophic Biological Risks](#), on ‘self-spreading vaccine’ technology, informed consent challenges of same, and ‘self-amplifying mRNA vaccines.’
- 2019/02/11 - President Trump signed [Executive Order 13859](#): *Maintaining American Leadership in Artificial Intelligence*. Directed and prioritized federal agency collaboration with industry for AI research and development.
- 2019/05/22 - [Congressional Research Service Opinion: An Overview of State and Federal Authority to Impose Vaccination Requirements](#) by Wen W. Shen
- 2019/06/11 - President Trump signed Executive [Order](#) 13874: *Modernizing the Regulatory Framework for Agricultural Biotechnology Products*. [84 Federal Register 27899](#).
- 2019/06/24 - Congress and President Trump passed Pandemic and All-Hazards Preparedness and Advancing Innovation Act - PL 116-22, [133 Stat. 905](#). Amended Public [Health](#) Service Act (42 U.S.C. 201), further consolidating federal power in HHS Secretary’s hands during public [health](#) emergencies, further merging public [health](#) and law enforcement systems, and further subordinating state, tribal, county and municipal governments and American civilians to direct federal control.
- 2019/09/19 - President Trump signed [Executive Order 13887](#): *Modernizing Influenza Vaccines in the United States to Promote National Security and Public [Health](#)*. Directed and prioritized federal agency collaboration with industry for rapid-deployment mRNA/DNA/LNP/nanotech bioweapon platforms misclassified as public [health](#) protection.
- 2019/10/04 - 10/19 - Roman Catholic Pope Francis hosted pagan Pachamama/Gaia worship ceremony in Vatican Garden, at Basilica of St. Peter, and Santa Maria

Traspontina [Church](#), and during Way of the Cross, until angry Catholics seized pagan statues and threw them into Tiber [River](#).

- 2019/10/18 - Johns Hopkins Center for [Health](#) Security conducted [Event 201](#):
“...a pandemic tabletop exercise that simulated a series of dramatic, scenario-based facilitated discussions, confronting difficult, true-to-life dilemmas associated with response to a hypothetical, but scientifically plausible, pandemic...”
 - 2019/12/12 - [Material Transfer Agreement](#) signed between US [Health](#) and Human Services (HHS) National Institutes of [Health](#) (NIH) National Institute for Allergies and [Infection](#) Diseases (NIAID), led by Anthony Fauci, [University](#) of North Carolina coronavirus researcher and patent-holder Ralph Baric, and Moderna, for “mRNA coronavirus vaccine candidates developed and jointly owned by NIAID and Moderna.”
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2020 - Present - Presidents Donald J. Trump, Joseph R. Biden

- 2020/01/27 - [US Secretary of Health and Human Services Determination that a Public Health Emergency Exists](#). Signed Jan. 31, 2020, effective Jan. 27, 2020. Renewed every 90 days since then, most recently Oct. 13, 2022. Also signed a ‘declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostics for detection and/or diagnosis of this novel coronavirus.’ The determination and declaration were recorded in the Federal Register as taking effect Feb. 4, 2020. [85 Federal Register 7316](#).
- 2020/01/30 - WHO Director-General Tedros Adhanom Ghebreyesus [declared Covid-19 outbreak a “public health emergency of international concern,”](#) (PHEIC) triggering the legal obligations of WHO member states under the 2005 International [Health](#) Regulations, to suspend national sovereignty and constitutional rights of citizens using the implementing domestic statutes and regulations they had adopted in compliance with the WHO IHR.
- 2020/02/04 - [US Secretary of Health and Human Services Declaration Under the Public Readiness and Emergency Preparedness Act for Medical Countermeasures Against COVID-19](#). 85 Federal Register 15198 (6 pages). Issued March 10, 2020, effective Feb. 4, 2020. Deployment of the domestic bioterrorism program against all American citizens under Covid-19 pretext.
- 2020/03/01 - HHS Centers for Medicare and Medicaid Services (CMS) [COVID-19 Emergency Declaration Blanket Waivers for Health Care Providers](#). Exempted [health](#) care providers from patient care standards and regulations that would legally apply in non-pandemic circumstances; authorized stripping patients of their rights

to have family members and pastors/rabbis visit them and advocate for them in the [hospital](#) or nursing home; supported [hospital](#) demands that law enforcement officers remove family and pastors from the premises by force; created conditions for [death protocols](#) of restraint, withheld [water](#) and nutrition, forcible administration of Remdesivir and forcible connection to ventilators under the ICD-10 codes.

- 2020/03/06 - Congress and President Trump passed Coronavirus Preparedness and Response Supplemental Appropriations Act - PL 116-123, [134 Stat. 146](#). \$8.3 billion to [Health](#) and Human Services, Centers for Disease Control and Prevention, National Institute of [Health](#), National Institute of Allergy and Infectious Diseases, Food and Drug Administration, Small Business Administration, Department of State and US Agency for International Development, for research and development of vaccines, therapeutics and diagnostics and other Covid programs.
- 2020/03/13 - [PanCAP Adapted U.S. Government Covid-19 Response Plan](#).
- 2020/03/13 - President Trump issued a [Stafford Act declaration](#) under the 1988 Stafford Act, and signed Proclamation 9994, *Declaring a National Emergency Concerning the Novel Coronavirus Disease (COVID-19) Outbreak*, under the 1975 National Emergencies Act. Renewed every year since, most recently by Biden in Feb. 2022. [85 Federal Register 15337](#).
- 2020/03/18 - Congress and President Trump passed Families First Coronavirus Response Act - PL 116-127, [134 Stat. 178](#). \$3.5 billion for Covid mass testing, supplemental nutrition (Department of Agriculture), sick leave, family medical leave, and unemployment compensation (Department of Labor) programs.
- 2020/03/24 - HHS Secretary Alex Azar issued Declaration of Emergency Use Authorization, declaring “that circumstances exist justifying the authorization of emergency use of medical devices, including alternative products used as medical devices.” [85 Federal Register 17335](#).
- 2020/03/27 - Congress and President Trump passed Coronavirus Aid, Relief, and Economic Security (CARES) Act - PL 116-136, [134 Stat. 281](#). 15 USC 9001. \$2.2 trillion in corporate and small business loans, household support and unemployment insurance, tax deferrals, aid to state and local governments, aid to universities and colleges, aid to K-12 schools, aid to hospitals and veterans programs, airline loans and grants, and \$10 billion for “Operation Warp Speed.”
- 2020/04/24 - Congress and President Trump passed Paycheck Protection Program and [Health](#) Care Enhancement Act - PL 116-139, [134 Stat. 620](#). \$75,000,000,000 for Public [Health](#) and Social Services Emergency Fund (first funded in 2005), “to remain available until expended, to prevent, prepare for, and respond to coronavirus, domestically or internationally” plus \$25,000,000,000 for research, development and deployment of Covid-19 tests.

- 2020/05/19 - [Advisory Opinion on the PREP Act and the March 10, 2020 Declaration Under the Act, April 17, 2020, as modified on May 19, 2020](#), by Robert P. Charrow of HHS [Office](#) of General Counsel. Legal opinion on statutory liability shields.
- 2020/05/29 - Supreme Court ruled in [South Bay United Pentecostal Church v. Newsom, 590 US ___, \(2020\)](#), denying role for federal judiciary in Constitutional review of executive and legislative acts taken during declared public [health](#) emergencies. Semi-reversed on rehearing, February 2021.
- 2020/07/20 - DOD-Pfizer [Base Agreement](#), through Advanced Technology International; 2020/07/21 - DOD-Pfizer [Statement of Work](#), through Advanced Technology International. Pfizer later argued (04/22/2022, *Jackson v. Ventavia*, [Motion to Dismiss](#)) that "Because of pandemic-related exigencies, the agreement was not a standard federal procurement contract, but rather a 'prototype' agreement executed pursuant to 10 U.S.C. § 2371b[.]...The [contract's Statement of Work] describes a 'large scale vaccine manufacturing demonstration' that imposes **no requirements relating to Good Clinical Practices ('GCP') or related FDA regulations.**" [10 USC Section 2371 renumbered 10 USC 4022](#), 01/01/2021
- 2020/08/26 - [HHS CDC Advisory Committee on Immunization Practices Meeting Summary Report](#). At p. 56 - "Dr. Cohn reminded everyone that under an EUA, vaccines are not allowed to be mandatory. Therefore, early in the vaccination phase individuals will have to be consented and cannot be mandated to be vaccinated." [Attorney Johnsen cited this interpretation of Section 564 in a footnote on p. 7 of her 07/06/2021 slip opinion, immediately citing the [judge's](#) 06/12/2021 [order](#) in *Bridges v. Houston Methodist* as "summarily rejecting" the argument.]
- 2020/12/27 - Consolidated Appropriations Act - PL 116-260, [134 Stat. 1182](#). \$2.3 trillion spending [bill](#), including \$900 billion for Covid programs.
- 2021/01/05 - Orange Book Transparency Act - PL 116-290, [134 Stat. 4889](#). Amendments to patent law under Federal Food Drug and Cosmetics Act, (21 USC 9)
- 2021/01/21 - HHS Secretary Norris Cochrane [notifies state governors that federal government will give 60 days notice](#) before terminating the "determination that a public [health](#) emergency exists" first issued by HHS Secretary Alex Azar effective 01/27/2020.
- 2021/03/11 - Congress and President Biden passed American Rescue Plan/Consolidated Appropriations Act. PL 117-2, [135 Stat. 4](#). Section 1401, Covid-19 Consumer Protection Act. Criminalized advocacy of alternative treatments under Federal Trade Commission provisions.
- 2021/04/02 - [Congressional Research Service Opinion: State and Federal Authority to Mandate COVID-19 Vaccination](#) (Version 1) by Wen W. Shen
- 2021/06/12 - Texas federal [judge](#) ruled in [Bridges v. Houston Methodist Hospital, 543 F. Supp. 3d 525](#) (S.D. Tex. 2021), finding that informed consent doesn't apply

to [hospital](#) workers, because the injections are government-authorized under FDA Emergency Use Authorization, therefore not part of experimental clinical trials or ordinary medical treatments, therefore [hospital](#) employees cannot be legally construed as human subjects or ordinary patients, therefore they have no individual, Constitutional liberties; rights to privacy and against government violation of bodily integrity; or rights to be secure in their persons against warrantless search and seizure.

- 2021/06/25 - [FDA EUA Pfizer Fact Sheet](#) addressing “option to accept or refuse.” This is only one of many versions issued between December 2020 and present; it’s the one cited by Attorney Johnsen in her legal opinion.
- 2021/07/06 - Dawn Johnsen, Deputy Attorney General, published [DOJ Opinion: Whether Section 564 of the Food, Drug, and Cosmetic Act Prohibits Entities from Requiring the Use of a Vaccine Subject to an Emergency Use Authorization](#). Related federal government’s position on legal status and regulatory control differences between Emergency Use Authorization (EUA) products, Investigational New Drugs (IND) and Investigational Device Exemptions (IDE).
- 2021/07/16 - Roman Catholic Pope Francis issued [Traditionis custodes](#), attempting to abrogate Pope Benedict’s 2007 *Summorum Pontificum*, and revoke the right of Catholic priests and faithful to celebrate the pre-1962, Traditional Latin Mass.
- 2021/07/29 - President Biden [directed](#) Department of Defense to “look into how and when they will add COVID-19 vaccination to the list of required vaccinations for members of the military.”
- 2021/08/24 - Department of Defense [order](#) from Secretary of Defense Lloyd [Austin](#), vaxx mandate on military personnel in Army, Navy, Air Force, Marines and Coast Guard.
- 2021/09 - HHS [FDA Guidance: Real-World Data - Assessing Electronic Health Records and Medical Claims Data To Support Regulatory Decision-Making for Drug and Biological Products](#)
- 2021/09/09 - President Biden signed Executive [Order](#) 14042, vaxx mandate on federal contractors. [86 Federal Register 50985](#).
- 2021/09/09 - President Biden signed Executive [Order](#) 14043, vaxx mandate on federal employees. [86 Federal Register 50989](#).
- 2021/09/09 - President Biden issued directive to Department of Labor Occupational Safety and [Health](#) Administration (OSHA), vaxx mandate on private employers with more than 100 employees.
- 2021/09/17 - President Biden signed Executive [Order](#) 14047, adding measles to the list of quarantinable communicable diseases authorizing HHS Secretary to use force to apprehend and detain people under [42 USC 264\(b\)](#) and [42 CFR 70.6](#). [86 Federal Register 52591](#).
- 2021/11 - HHS [FDA Guidance: Real-World Data - Assessing Registries to Support Regulatory Decision-Making for Drug and Biological Products](#)

- 2021/11/05 - President Biden issued directive to Department of [Health](#) and Human Services Center for Medicare and Medicaid Services (CMS), vaxx mandate on [health](#) care workers at hospitals, nursing homes and other federally-funded facilities.
- 2021/11/17 - HHS Interim Final Rule - Possession, Use, and Transfer of Select Agents and Toxins—Addition of SARS-CoV/SARS-CoV-2 Chimeric Viruses Resulting From Any Deliberate Manipulation of SARS-CoV-2 To Incorporate Nucleic Acids Coding for SARS-CoV Virulence Factors to the HHS List of Select Agents and Toxins. [86 Federal Register 64075](#). Chimeric, lab-weaponized SARS-CoV-2 added to list of agents that “have the potential to pose a severe threat to public [health](#) and safety” under 42 CFR 73.3. Attempt to block accountability by preemptively reclassifying bioweapons as legally identical to pandemics, to block international law claims brought under the theory that SARS-CoV-2 is a bioweapon, and not a pandemic. If classified as a bioweapon, the Public [Health](#) Emergency of International Concern (international) and public [health](#) emergency (federal) legal frameworks would be nullified, instead bringing to [bear](#) federal and international laws prohibiting chemical and biological weapons.
- 2021/12/02 - HHS Final Rule - National Vaccine Injury Compensation Program: Adding the Category of Vaccines Recommended for Pregnant Women to the Vaccine Injury Table - [86 Federal Register 68423](#). Added vaccines recommended for pregnant women to the list of vaccines subject to the 1986 VICP compensation scheme, so as add another hurdle to civil suits against Covid-19 injection manufacturers, even though the products had not yet been added to the childhood vaccine schedule that otherwise governs access to VICP scheme. Because CDC does recommend them for pregnant women.
- 2021/12/27 - Congress and President Biden passed National Defense Authorization Act FY2022 - PL 117-81, [135 Stat. 1541](#). At Section 716, established military vaxx tracking system, including refusals, under 10 USC 1110 (originally re anthrax vaxx). At Section 6501, authorized US government to engage with [Bill](#) Gates Coalition for Epidemic Preparedness Innovations (CEPI). [More coverage](#).
- 2022/01/13 - Supreme Court ruled in [Missouri v. Biden \(21 A 240\)](#), [Louisiana v. Biden \(21 A. 241\)](#), [595 US ___, \(2022\)](#), asserting federal funding for hospitals and nursing homes voids Constitutional protection for employees individual bodily integrity and informed consent to medical treatment.
- 2022/02/07 - [Congressional Research Service Opinion: State and Federal Authority to Mandate COVID-19 Vaccination](#)
- 2022/02/10 - Supreme Court leaked draft opinion in Dobbs v. Jackson Women's [Health](#), [leaked draft opinion](#) by Justice Samuel Alito. SCOTUS poised to explicitly deny the principle of Constitutionally-protected inalienable individual rights to personal privacy, conscience, bodily integrity, or liberty, against State exercise of authority. Final ruling issued 06/24/2022.

- 2022/03/09 - President Biden signed Executive [Order](#) 14067, *Ensuring Responsible Development of Digital Assets*, on Central Bank Digital Currencies)
- 2022/03/15 - Congress and President Biden passed Consolidated Appropriations Act - PL 117-103, [136 Stat. 49](#). \$1,274,678,000 for the Public [Health](#) and Social Services Emergency Fund (HHS slush fund established in 2005). \$780,000,000 for new domestic bioweapons production, classified as 'security countermeasures' under the Public [Health](#) Service Act as amended by 2004 Project Bioshield Act, [42 USC 247d-6b\(c\)\(1\)\(B\)](#); \$845,000,000 to stock the Strategic National Stockpile established 1998, controlled by the CDC within HHS [42 USC 247d-6b\(a\)](#); \$300,000,000 "to prepare for or respond to an influenza pandemic," including federally-funded construction or renovation of privately-owned pharmaceutical manufacturing facilities, if the Secretary of [Health](#) and Human Services finds such construction or renovation necessary; \$1,000,000,000 to establish ARPA-H: Advanced Research Program Agency - [Health](#), to conduct research and development of bioweapons misbranded as public [health](#) measures; \$3,880,000,000 to US Agency for International Development (US-AID) for programs mislabeled as 'Global [Health](#) Programs,' including immunization programs, HIV/AIDS programs, The GAVI Alliance [population-control zealot [Bill](#) Gates' Global Alliance for Vaccines and Immunization] and a multilateral vaccine development partnership, for, among other projects, "experimental contraceptive drugs, devices and medical procedures."
- 2022/05/17 - [Congressional Research Service Opinion: State and Federal Authority to Mandate COVID-19 Vaccination](#). (Version 9)
- 2022/05/17 - [Congressional Research Service Opinion: Status of Federal COVID-19 Vaccination Mandate Litigation](#). (Version 7)
- 2022/06/24 - Dobbs v. Jackson Womens [Health](#) SCOTUS decision released.
- 2022/07/15 - HHS Secretary Xavier Becerra extended 01/27/2020 [determination that 'public health emergency' exists](#).
- 2022/07/22 - HHS Secretary Xavier Becerra elevated Administration for Strategic Preparedness and Response (ASPR) from staff division to operating division, still under HHS Assistant Secretary Dawn O'Connell.
- 2022/09/12 - President Biden signed Executive [Order](#) 14081 - *Advancing Biotechnology and Biomanufacturing Innovation for a Sustainable, Safe, and Secure American Bioeconomy*. [87 Federal Register 56849](#).
- 2022/09/28 - HHS-FDA Proposed Rules: *Protection of Human Subjects and Institutional Review Boards*. [87 Federal Register 58733](#)
- 2022/10/13 - HHS Secretary Xavier Becerra extended 01/27/2020 [determination that 'public health emergency' exists](#).
- 2022/10/13 - [Boston University](#) researchers, funded by Fauci's NIAID, publish preprint paper on their gain-of-function/DURC research combining the toxicity of the original Wuhan strain of SARS-CoV-2 with the increased transmissibility of the Omicron variant to

achieve 80% mortality in transgenic, humanized mice expressing the ACE-2 receptor. [Role of spike in the pathogenic and antigenic behavior of SARS-CoV-2 BA.1 Omicron](#)

- 2022/10/18 - President Biden [National Security Memorandum \(NSM-15\) on Countering Biological Threats, Enhancing Pandemic Preparedness, and Achieving Global Health Security](#) and [National Biodefense Strategy Implementation Plan](#).
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Summer 2022 - Pending Federal Legislation

- [National Defense Authorization Act](#) for FY2023 - Pending, HR7900. Section 6901 - Global [Health](#) Security Act. Authorizes, creates, funds globalized military-[health](#) structure, linking US military to global genocide apparatus operating under WHO frameworks.
- Global Pandemic Prevention and Biosecurity Act - Pending, [HR3424](#) and [S1737](#). Also included in NDAA for FY2023 at Section 6901.
- [2022 Covid Supplemental Appropriations Act](#) - Pending, HR7007. Authorizes \$10.6 billion for Covid bioweapon development and deployment, including “up to \$9,850,000,000 to Biomedical Advanced Research and Development Authority [BARDA, established 2006] for advanced research and development, manufacturing, production, and purchase, at the discretion of the Secretary of [Health](#) and Human Services, of vaccines, therapeutics, diagnostics, and supplies.”
- [2022 Research Investment to Spark the Economy \(RISE\) ACT](#) - Pending, S.289. Senate counterpart to Cures 2.0 Act/HR6000, Title V, Section 502. Authorizes billions in funding for the Departments of Agriculture, Commerce, Defense, Education, Energy, the Interior, [Health](#) and Human Services, and Transportation, National Aeronautics and Space Administration (NASA), National [Science](#) Foundation, and Environmental Protection Agency to provide support for research regarding COVID-19 (i.e., coronavirus disease 2019) or research disrupted by the COVID-19 pandemic. Support may be used to provide supplemental funding to extend the duration of a grant...that was awarded prior to enactment, or to expand the purposes of such a grant; issue awards to research the effects of the current pandemic and potential future pandemics; and provide flexibility on awards to account for facility closures or other limitations during the COVID-19 public [health](#) emergency.
- [2022 PASTEUR Act](#) - Pending, HR 3932. (41 pages). Pioneering Anti-microbial Subscriptions To End Upsurging Resistance Act. Would create subscription-based procurement contracts between the US government and pharmaceutical corporations for ongoing, open-ended development, purchase and deployment of drugs alleged to treat

antibiotic-resistant infections. Appropriates \$11 billion for program. Program to be developed by committee comprised of National Institute of Allergy and Infectious Diseases, Centers for Disease Control and Prevention, Biomedical Advanced Research and Development Authority, Food and Drug Administration, Centers for Medicare & Medicaid Services, Veterans [Health](#) Administration, and Department of Defense.

- [2022 Cures 2.0 Act](#) - Pending, HR6000. (173 pages.) Would legally establish Covid-[infection](#) injury and Covid-19 bioweapon injection injury as “long Covid,” (erasing injection-caused injury as a separate diagnostic classification) and appropriate research and treatment funding; would establish genomic testing program for children and teens (corroborating evidence that government developed the bioweapons to cause listed harms and anticipates observing those effects in the population); would establish pharmacogenetic consulting and other programs. Title V, Section 502 is [House](#) counterpart to S.289, RISE Act (see above), to authorize billions in funding for the Departments of Agriculture, Commerce, Defense, Education, Energy, the Interior, [Health](#) and Human Services, and Transportation, National Aeronautics and Space Administration (NASA), National [Science](#) Foundation, and Environmental Protection Agency to provide support for research regarding COVID-19 (i.e., coronavirus disease 2019) or research disrupted by the COVID-19 pandemic.

COVID-19 injectable bioweapons as case study in legalized, government-operated domestic bioterrorism. Or: why there won't be any civil suits, or compensatory damages for injured victims or survivors of dead victims.

Since first realizing the implications of the many Congressional statutes and [Health](#) and Human Services regulations adopted to create and operate the bioterrorism program, mostly between 1997 and the present, I've been intermittently finding the specific citations for each statement while researching related issues.

Some statements are simply logical deductions from the first premise, corroborated by the observable actions and inactions of Food and Drug Administration officials as the observable injuries and deaths mount up in the American people.

Others are specifically written into the laws, but I don't yet have the citations because I've prioritized my research time investigating other issues related to the bioterrorism program.

I'm posting the information as I understand it today [June 9, 2022], despite those limitations, in case it's useful for readers who also follow FDA Vaccine and Related Biological Products Advisory Committee (VRBPAC) reporting by [Toby Rogers](#), [Igor Chudov](#), [Steve Kirsch](#), [Jessica Rose](#), and others.

They continue to rightly raise public awareness and alarm about FDA's ongoing failure to protect the public from the Emergency Use Authorized (EUA) products.

But they don't address the main reason **why** FDA is acting as it is.

FDA is not pulling the EUA products from the market or stopping the 'vaccination' campaign because [Health](#) and Human Services Secretary Xavier Becerra and FDA Commissioner Robert Califf are running the US government's bioterrorism program jointly with Defense Secretary Lloyd [Austin](#), Department of Justice Attorney General Merrick Garland, Department of Homeland Security Secretary Alejandro Mayorkas, Pfizer CEO Albert Bourla, Moderna CEO Stéphane Bancel, and World [Health](#) Organization Director-General Tedros Adhanom Ghebreyesus.

Main Premise

Use of EUA-covered medical countermeasure (MCM) products including masks, PCR tests, mRNA and DNA injections, and other drugs, devices and biologics, once designated as such by the Secretary of [Health](#) and Human Services ([March 10, 2020, retroactive to February 4, 2020](#)) "**shall not be considered to constitute a clinical investigation.**" 21 USC 360bbb-3(k). FDA EUA law, adopted 1997 and amended 2003, 2004, 2005, 2013, 2017.

This is true no matter how untested, unmonitored, unsafe, or ineffective they are, no matter whether their harmfulness to human [health](#) and uselessness for [infection](#)-control are known before use, or discovered afterward.

Legal implications derived from the main premise:

1. **There is no stopping condition.**
2. EUA products are exempt from laws regulating researcher use of investigational, experimental drugs, devices and biologics on human beings.
3. EUA products are exempt from laws regulating physician use of approved drugs, devices and biologics as medical treatments for patients.
4. There are no manufacturers of experimental products (EUA products are not part of any clinical investigation, and therefore not experimental.)
5. There are no government or private contracts for purchase of experimental products; there are only contracts for '[large scale vaccine manufacturing demonstrations](#).'
6. There is no act of administration of any experimental products.
7. There are no nurses or pharmacists administering experimental products.
8. There are no human subjects (of experiments) or patients (of physicians providing treatment) receiving experimental products: no victims.
9. There is no party responsible for the wellbeing of recipients after administration of EUA products.
10. There is no treatment group and no control group.
11. Human beings administering EUA products have no **informed consent** obligations to provide information about ingredients, risks, benefits, alternatives, or the option to accept

or refuse the products. See 21 USC 360bbb-3(e)(1)(A)(ii)) waiving informed consent for unapproved products (2004); 21 USC 360bbb-3(e)(2)(A) waiving informed consent for unapproved use of an approved product (2004); 21 USC 355(i)(4) waiving informed consent for experimental products classified by HHS as ‘minimal risk’ drugs (2016); 21 USC 360j(g)(3)(D)(i) waiving informed consent for experimental ‘minimal risk’ devices (2016).

12. Human beings receiving EUA products have no **informed consent** rights to receive information about ingredients, risks, benefits, alternatives, or the option to accept or refuse the products. See citations, bullet point above.
13. There are no Institutional Review Boards supervising administration of the experimental products.
14. There are no safety standards for EUA products.
15. There are no efficacy standard for EUA products. See 21 USC 360bbb-3(c)(2)(A), 1997, 2003, 2004, re: ‘may be effective.’
16. There are no clinical investigators studying the effects of EUA products on human subjects.
17. There are no doctors, nurses, or other treatment providers providing experimental treatment to their patients subject to the Hippocratic Oath (“first do no harm”) using EUA products.
18. There is no coordinated, public, federal government monitoring of recipients after receiving the products for adverse effects and deaths.
19. There is no coordinated, public, federal government data collection or analysis.
20. There is no legal requirement for medical supervision during product administration.
21. There is no legal requirement for recipient monitoring after product administration.
22. ‘Real world evidence’ — mass administration of products to general public, followed by collection of private/proprietary information about the effects, from [health](#) insurance systems, government databases ([Medicare](#), Medicaid, Defense Medical Epidemiology Database, Veterans [Health](#) Administration) and other private databases — is authorized for the purposes of FDA regulatory decisions. See 21 USC 355g. 2016.
23. There is no requirement for individual prescriptions to be written prior to dispensing EUA products, and products dispensed without prescriptions “shall not be deemed adulterated or misbranded.” See 21 USC 360bbb-3a(d). 2013.
24. Manufacturers, as contractors, are considered HHS employees for purposes of legal immunity under Federal Tort Claims Act. See 42 USC 247d-6a(d)(2)(A).
25. DOD is authorized to contract with pharmaceutical corporations to conduct ‘prototype’ experiments on the general public, and under such contracts, is exempt from legal obligation to comply with Good Clinical Practices or other FDA regulations. See 10 USC 2371b (2015), renumbered 10 USC 4022 (Jan. 1, 2021, effective Jan. 1, 2022)

26. One of the factors to be considered by HHS secretary in making determinations about EUA products (qualified security countermeasures) and use of Special Reserve Fund/Strategic National Stockpile appropriations to procure them is "whether there is a lack of a significant commercial market for the product at the time of procurement, other than as a security countermeasure." See 42 USC 247d-6b (c)(5)(B)(iii)
27. There are no required standards for quality-control in manufacturing; no inspections of manufacturing procedures; no prohibition on wide variability among lots; no prohibition on adulteration; and no required compliance with Current Good Manufacturing Practices. EUA products, even though unregulated and non-standardized, "shall not be deemed adulterated or misbranded." See 21 USC 360bbb-3a(c). 2013.
28. There are no labeling requirements regarding the contents or ingredients in EUA products. 21 USC 360bbb-3(e)(2)(B)(ii). 2004.
29. There is no limitation of administration of EUA products past their expiration dates.
30. There cannot be clinical trial fraud, because there are no clinical investigations, no investigational drugs, no investigators and no human subjects.
31. There are no marketing standards.
32. There cannot be consumer fraud, because the only legal parties to the financial transactions are the US government (DOD) as buyer; the US government (HHS) as regulator authorizing exemptions from consumer protection laws that otherwise apply to medical products; and the pharmaceutical corporations as sellers, contracted to develop and manufacture the products. There are no commercial pharmaceutical products, no commercial marketplace, and no commercial market consumers.
33. There is no access to courts for judicial review of the facts or law relating to HHS Secretary declarations of EUA products, which are committed to agency discretion. See 42 USC 247d-6d(b)(7). 2005.
34. There is no access for plaintiffs, to civil courts for judicial review, and no entity to whom civil liability can attach, for injuries and deaths caused by declared covered countermeasures, unless and until FDA/HHS and/or Attorney General/DOJ file enforcement action against manufacturers and prove willful misconduct proximate to injury or death, but HHS and DOJ have operated the EUA product program together with the manufacturers since inception, and will not prosecute their co-conspirators. See 42 USC 247d-6d. 2005.
35. Even if there were access to courts for judicial review, and a fact-finder found evidence of harms caused by administration of products to recipients, and even evidence that those who caused the harms, by developing, manufacturing, distributing and/or administering the EUA products, knew the EUA products were toxic and knew their own actions were harmful, "just following orders" is an authorized, legal defense. See 42 USC 247d-6d(c)(4). 2005.

Summary:

There are no actions that can be legally classified as crimes or civil torts; there are no medical battery or homicide victims, or plaintiffs; and there are no medical batterers or murderers. Because legally, nothing has been done, and no one has done anything, to anyone else. The recursive loop can be infinite, as covered countermeasures are developed, authorized and deployed, through HHS Secretary EUA declarations, as treatments for complications from prior countermeasures.

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